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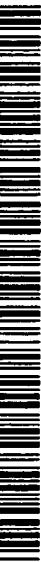
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(54) Title: TRANSGENIC PLANTS EXPRESSING PHOTORHABDUS TOXIN

(57) Abstract: Novel polynucleotide sequences that encode insect toxins TcdA and TcbA have base compositions that differ substantially from the native genes, making them more similar to plant genes. The new sequences are suitable for use for high expression in both monocots and dicots. Transgenic plants with a genome comprising a nucleic acid of SEQ ID NO: 3 or SEQ ID NO:4 are insect resistant.

TRANSGENIC PLANTS EXPRESSING PHOTORHABDUS TOXIN

BACKGROUND OF THE INVENTION

As reported in WO98/08932, protein toxins from the genus *Photorhabdus* have been shown to have oral toxicity against insects. The toxin complex produced by *Photorhabdus luminescens* (W-14), for example, has been shown to contain ten to fourteen proteins, and it is known that these are produced by expression of genes from four distinct genomic regions: *tca*, *tcb*, *tcc*, and *tcd*.
10 WO98/08932 discloses nucleotide sequences for the native toxin genes.

Of the separate toxins isolated from *Photorhabdus luminescens* (W-14), those designated Toxin A and Toxin B are especially potent against target insect species of interest, for example corn rootworm. Toxin A is comprised of two different subunits. The native gene *tcdA* (SEQ ID NO:1) encodes protoxin TcdA (see SEQ ID NO:1). As determined by mass spectrometry, TcdA is processed by one or more proteases to provide Toxin A.
20 More specifically, TcdA is an approximately 282.9 kDa protein (2516 aa) that is processed to provide TcdAii, an approximately 208.2 kDa (1849 aa) protein encoded by nucleotides 265-5811 of SEQ ID NO:1, and TcdAiii, an approximately 63.5 kDa (579 aa) protein encoded by nucleotides 5812-7551 of SEQ ID NO:1.
25

Toxin B is similarly comprised of two different subunits. The native gene *tcbA* (SEQ ID NO:2) encodes protoxin TcbA (see SEQ ID NO:2). As determined by mass spectrometry, TcbA is processed by one or more proteases to provide Toxin B. More specifically, TcbA is an approximately 280.6 kDa (2504 aa) protein that is processed to provide TcbAii, an approximately 207.7 kDa (1844 aa) protein encoded by nucleotides 262-5793 of SEQ ID NO:2 and TcbAiii, an approximately 62.9 kDa (573 aa) protein encoded by nucleotides 5794-7512 of SEQ ID NO:2.
35

The native *tcdA* and *tcbA* genes are not well suited for high level expression in plants. They encode multiple destabilization sequences, mRNA splice sites, polyA addition sites and other possibly detrimental sequence motifs. In addition, the codon compositions are not like those of plant genes. WO98/08932 gives general guidance on how the toxin genes could be reengineered to more efficiently expressed in the cytoplasm of plants, and describes how plants can be transformed to incorporate the *Photorhabdus* toxin genes into their genomes.

SUMMARY OF THE INVENTION

In a preferred embodiment, the invention provides novel polynucleotide sequences that encode TcdA and TcbA. The novel sequences have base compositions that differ substantially from the native genes, making them more similar to plant genes. The new sequences are suitable for use for high expression in both monocots and dicots, and this feature is designated by referring to the sequences as the "hemicot" criteria, which is set forth in detail hereinafter. Other important features of the sequences are that potentially deleterious sequences have been eliminated, and unique restriction sites have been built in to enable adding or changing expression elements, organellar targeting signals, engineered protease sites and the like, if desired.

In a particularly preferred embodiment, the invention provides polynucleotide sequences that satisfy hemicot criteria and that comprise a sequence encoding an endoplasmic reticulum signal or similar targeting sequence for a cellular organelle in combination with a sequence encoding TcdA or TdbA.

More broadly, the invention provides engineered nucleic acids encoding functional *Photorhabdus* toxins wherein the sequences satisfy hemicot criteria.

The invention also provides transgenic plants with genomes comprising a novel sequence of the invention that imparts functional activity against insects.

5

BRIEF DESCRIPTION OF SEQUENCES

SEQ ID NO:1 is the native *tcdA* DNA sequence together with the corresponding encoded amino acid sequence for *TcdA*.

SEQ ID NO:2 is the native *tcbA* DNA sequence together 10 with the corresponding encoded amino acid sequence for *TcbA*.

SEQ ID NO:3 is an artificial sequence encoding *TcdA* that is suitable for expression in monocot and dicot plants.

15 SEQ ID NO:4 is an artificial sequence encoding *TdbA* that is suitable for expression in monocot and dicot plants.

SEQ ID NO:5 is an artificial hemicot sequence that encodes the 21 amino acid ER signal peptide of 15 kDa 20 zein from Black Mexican Sweet maize.

SEQ ID NO:6 is an artificial hemicot sequence that encodes for the full-length native *TcdA* protein (amino acids 22-2537) fused to the modified 15 kDa zein endoplasmic reticulum signal peptide (amino acids 1-21).

25

DETAILED DESCRIPTION

The native *Photorhabdus* toxins are protein complexes that are produced and secreted by growing bacteria cells of the genus *Photorhabdus*. Of particular interest are the proteins produced by the species *Photorhabdus* 30 *luminescens*. The protein complexes have a molecular size of approximately 1,000 kDa and can be separated by SDS-PAGE gel analysis into numerous component proteins. The toxins contain no hemolysin, lipase, type C phospholipase, or nuclease activities. The toxins 35 exhibit significant toxicity upon ingestion by a number of insects.

A unique feature of *Photorhabdus* is its bioluminescence. *Photorhabdus* may be isolated from a variety of sources. One such source is nematodes, more particularly nematodes of the genus *Heterorhabditis*.
5 Another such source is from human clinical samples from wounds, see Farmer et al. 1989 J. Clin. Microbiol. 27 pp. 1594-1600. These saprophytic strains are deposited in the American Type Culture Collection (Rockville, MD) ATCC #s 43948, 43949, 43950, 43951, and 43952, and are
10 incorporated herein by reference. It is possible that other sources could harbor *Photorhabdus* bacteria that produce insecticidal toxins. Such sources in the environment could be either terrestrial or aquatic based.

The genus *Photorhabdus* is taxonomically defined as a
15 member of the Family *Enterobacteriaceae*, although it has certain traits atypical of this family. For example, strains of this genus are nitrate reduction negative, yellow and red pigment producing and bioluminescent. This latter trait is otherwise unknown within the
20 *Enterobacteriaceae*. *Photorhabdus* has only recently been described as a genus separate from the *Xenorhabdus* (Boemare et al., 1993 Int. J. Syst. Bacteriol. 43, 249-255). This differentiation is based on DNA-DNA hybridization studies, phenotypic differences (e.g.,
25 presence (*Photorhabdus*) or absence (*Xenorhabdus*) of catalase and bioluminescence) and the Family of the nematode host (*Xenorhabdus*; *Steinernematidae*,
Photorhabdus; *Heterorhabditidae*). Comparative, cellular fatty-acid analyses (Janse et al. 1990, Lett. Appl. Microbiol 10, 131-135; Suzuki et al. 1990, J. Gen. Appl. Microbiol., 36, 393-401) support the separation of
30 *Photorhabdus* from *Xenorhabdus*.

Currently, the bacterial genus *Photorhabdus* is comprised of a single defined species, *Photorhabdus*
35 *luminescens* (ATCC Type strain #29999, Poinar et al., 1977, Nematologica 23, 97-102). A variety of related

strains have been described in the literature (e.g.,
Akhurst et al. 1988 J. Gen. Microbiol., 134, 1835-1845;
Boemare et al. 1993 Int. J. Syst. Bacteriol. 43 pp. 249-
255; Putz et al. 1990, Appl. Environ. Microbiol., 56,
5 181-186).

The following toxin producing *Photorhabdus* strains
have been deposited:

strain	accession number	date of deposit
W-14	ATCC 55397	March 5, 1993
WX1	NRRL B-21710	April 29, 1997
WX2	NRRL B-21711	April 29, 1997
WX3	NRRL B-21712	April 29, 1997
WX4	NRRL B-21713	April 29, 1997
WX5	NRRL B-21714	April 29, 1997
WX6	NRRL B-21715	April 29, 1997
WX7	NRRL B-21716	April 29, 1997
WX8	NRRL B-21717	April 29, 1997
WX9	NRRL B-21718	April 29, 1997
WX10	NRRL B-21719	April 29, 1997
WX11	NRRL B-21720	April 29, 1997
WX12	NRRL B-21721	April 29, 1997
WX14	NRRL B-21722	April 29, 1997
WX15	NRRL B-21723	April 29, 1997
H9	NRRL B-21727	April 29, 1997
Hb	NRRL B-21726	April 29, 1997
Hm	NRRL B-21725	April 29, 1997
HP88	NRRL B-21724	April 29, 1997
NC-1	NRRL B-21728	April 29, 1997
W30	NRRL B-21729	April 29, 1997
WIR	NRRL B-21730	April 29, 1997
B2	NRRL B-21731	April 29, 1997
ATCC 43948	ATCC 55878	November 5, 1996
ATCC 43949	ATCC 55879	November 5, 1996
ATCC 43950	ATCC 55880	November 5, 1996
ATCC 53951	ATCC 55881	November 5, 1996
ATCC 43952	ATCC 55882	November 5, 1996
DEPI	NRRL B-21707	April 29, 1997
DEP2	NRRL B-21708	April 29, 1997
DEP3	NRRL B-21709	April 29, 1997
P. zealandrica	NRRL B-21683	April 29, 1997
P. hepialus	NRRL B-21684	April 29, 1997
HB-Arg	NRRL B-21685	April 29, 1997
HB Oswego	NRRL B-21686	April 29, 1997
Hb Lewiston	NRRL B-21687	April 29, 1997
K-122	NRRL B-21688	April 29, 1997
HMGD	NRRL B-21689	April 29, 1997
Indicus	NRRL B-21690	April 29, 1997
GD	NRRL B-21691	April 29, 1997
PWH-5	NRRL B-21692	April 29, 1997
Megidis	NRRL B-21693	April 29, 1997
HF-85	NRRL B-21694	April 29, 1997
A. Cows	NRRL B-21695	April 29, 1997
MP1	NRRL B-21696	April 29, 1997
MP2	NRRL B-21697	April 29, 1997
MP3	NRRL B-21698	April 29, 1997
MP4	NRRL B-21699	April 29, 1997
MP5	NRRL B-21700	April 29, 1997
GL98	NRRL B-21701	April 29, 1997
GL101	NRRL B-21702	April 29, 1997
GL138	NRRL B-21703	April 29, 1997
GL155	NRRL B-21704	April 29, 1997
GL217	NRRL B-21705	April 29, 1997
GL257	NRRL B-21706	April 29, 1997

All strains were deposited in accordance with the terms of the Budapest Treaty. Strains having

accession numbers prefaced by "ATTC" were deposited on the indicated date in the American Type Culture Collection, 12301 Parklawn Drive, Rockville, MD 20852 USA. Strains prefaced by "NRRL" were

5 deposited on the indicated date in the Agricultural Research Service Patent Culture Collection (NRRL), National Center for Agricultural Utilization Research, ARS-USDA, 1815 North University St., Peoria IL 61604 USA.

10 The present invention provides hemicot nucleic acid sequences encoding toxins from any *Photorhabdus* species or strain that produces a toxin having functional activity. Hemicot nucleic acid sequences encoding proteins homologous to such toxins are also encompassed 15 by the invention.

Several terms that are used herein have a particular meaning and are defined as follows:

By "functional activity" it is meant herein that the protein toxins) function as insect control agents in that 20 the proteins are orally active, or have a toxic effect, or are able to disrupt or deter feeding, which may or may not cause death of the insect. When an insect comes into contact with an effective amount of toxin delivered via transgenic plant expression, formulated protein 25 compositions), sprayable protein compositions), a bait matrix or other delivery system, the results are typically death of the insect, or the insects do not feed upon the source which makes the toxins available to the insects.

30 By "homolog" it is meant an amino acid sequence that is identified as possessing homology to a reference *Photorhabdus* toxin polypeptide amino acid sequence.

By "homology" it is meant an amino acid sequence that has a similarity index of at least 33% and/or an 35 identity index of at least 26% to a reference *Photorhabdus* toxin polypeptide amino acid sequence, as

scored by the GAP algorithm using the B10sum 62 protein scoring matrix Wisconsin Package Version 9.0, Genetics Computer Group GCG), Madison, WI).

By "identity" is meant an amino acid sequence that 5 contains an identical residue at a given position, following alignment with a reference *Photorhabdus* toxin polypeptide amino acid sequence by the GAP algorithm.

By the use of the term "Photorhabdus toxin" it is meant any protein produced by a *Photorhabdus* 10 microorganism strain which has functional activity against insects, where the *Photorhabdus* toxin could be formulated as a sprayable composition, expressed by a transgenic plant, formulated as a bait matrix, delivered via baculovirus, or delivered by any other applicable 15 host or delivery system.

By the use of the term "toxic" or "toxicity" as used herein it is meant that the toxins produced by *Photorhabdus* have "functional activity" as defined herein.

By "substantial sequence homology" is meant either: 20 a DNA fragment having a nucleotide sequence sufficiently similar to another DNA fragment to produce a protein having similar biochemical properties; or a polypeptide having an amino acid sequence sufficiently similar to 25 another polypeptide to exhibit similar biochemical properties.

As with other bacterial toxins, the rate of mutation of the bacteria in a population causes many related toxins slightly different in sequence to exist. Toxins 30 of interest here are those which produce protein complexes toxic to a variety of insects upon exposure, as described herein. Preferably, the toxins are active against *Lepidoptera*, *Coleoptera*, *Homoptera*, *Diptera*, *Hymenoptera*, *Dictyoptera* and *Acarina*. The inventions 35 herein are intended to capture the protein toxins homologous to protein toxins produced by the strains

herein and any derivative strains thereof, as well as any protein toxins produced by *Photorhabdus*. These homologous proteins may differ in sequence, but do not differ in function from those toxins described herein.

5 Homologous toxins are meant to include protein complexes of between 300 kDa to 2,000 kDa and are comprised of at least two 2) subunits, where a subunit is a peptide which may or may not be the same as the other subunit. Various protein subunits have been identified and are taught in
10 the Examples herein. Typically, the protein subunits are between about 18 kDa to about 230 kDa; between about 160 kDa to about 230 kDa; 100 kDa to 160 kDa; about 80 kDa to about 100 kDa; and about 50 kDa to about 80 kDa.

As discussed above, some *Photorhabdus* strains can be
15 isolated from nematodes. Some nematodes, elongated cylindrical parasitic worms of the phylum *Nematoda*, have evolved an ability to exploit insect larvae as a favored growth environment. The insect larvae provide a source of food for growing nematodes and an environment in which
20 to reproduce. One dramatic effect that follows invasion of larvae by certain nematodes is larval death. Larval death results from the presence of, in certain nematodes, bacteria that produce an insecticidal toxin which arrests larval growth and inhibits feeding activity.

25 Interestingly, it appears that each genus of insect parasitic nematode hosts a particular species of bacterium, uniquely adapted for symbiotic growth with that nematode. In the interim since this research was initiated, the name of the bacterial genus *Xenorhabdus*
30 was reclassified into the *Xenorhabdus* and the *Photorhabdus*. Bacteria of the genus *Photorhabdus* are characterized as being symbionts of *Heterorhabditus* nematodes while *Xenorhabdus* species are symbionts of the *Steinernema* species. This change in nomenclature is
35 reflected in this specification, but in no way should a

change in nomenclature alter the scope of the inventions described herein.

The peptides and genes that are disclosed herein are named according to the guidelines recently published in 5 the Journal of Bacteriology "Instructions to Authors" p. i-xii Jan. 1996), which is incorporated herein by reference.

Transformation methods useful in carrying out the invention are well known, and are described, for example, 10 in WO98/08932.

Hemicot tcdA and tcbA

SEQ ID NO: 3 is the nucleotide sequence for an engineered *tcdA* gene in accordance with the invention.

SEQ ID NO: 4 is the nucleotide sequence for an engineered 15 *tcbA* gene in accordance with the invention.

The following Tables 1 and 2 identify significant features of the engineered *tcdA* and *tcbA* genes.

Table 1
tcdA

Feature	nucleotides of SEQ ID NO:3
<i>NcoI</i>	1-6
<i>HindIII</i>	48-53
<i>KpnI</i>	246-254
sequence encoding <i>TcbAii</i>	267-5798
<i>NheI</i>	333-338
<i>BglII</i>	1215-1220
<i>ClaI</i>	2604-2609
<i>PstI</i>	4015-4020
<i>AgeI</i>	5088-5093
<i>MunI</i>	5598-5603
<i>XbaI</i>	5778-5783
sequence encoding <i>TcbAii</i>	5799-7517
<i>AflII</i>	5853-5858
<i>SphI</i>	6439-6444
<i>SfuI</i>	7392-7397
<i>SacI</i>	7519-7524
<i>XhoI</i>	7522-7527
<i>StuI</i>	7528-7533
<i>NotI</i>	7533-7538

20

Table 2
tcbA

Feature	nucleotides of SEQ ID NO:5
<i>NcoI</i>	1-6
<i>HindIII</i>	48-53

<i>Kpn</i> I	246-251
sequence encoding <i>TcbAii</i>	267-5798
<i>Nhe</i> I	333-338
<i>Bgl</i> II	1215-1220
<i>Cl</i> AI	2604-2609
<i>Pst</i> I	4015-4020
<i>Age</i> I	5088-5093
<i>Mun</i> I	5598-5603
<i>Xba</i> I	5778-5783
sequence encoding <i>TcbAiii</i>	5799-7517
<i>Afl</i> II	5853-5858
<i>Sph</i> I	6439-6444
<i>Sfu</i> I	7392-7397
<i>Sac</i> I	7519-7524
<i>Sfu</i> I	7392-7397
<i>Sac</i> I	7519-7524
<i>Xho</i> I	7522-7527
<i>Stu</i> I	7528-7533
<i>Not</i> I	7535-7540

It should be noted that the proteins encoded by the plant-optimized *tcdA* (SEQ ID NO:3) and *tcbA* (SEQ ID NO:5) differ from the native proteins by the addition of 5 an Ala residue at position #2. This modification was made to accommodate the *Nco*I site which spans the ATG start codon.

10 The following Table 3 compares the codon composition of the engineered *tcdA* gene of SEQ ID NO:3 and engineered *tcbA* gene of SEQ ID NO:5 with the codon compositions of the native genes, the typical dicot genes, and maize genes.

Table 3

amino acid	codon	% in SEQ ID NO:3	% in <i>tcdA</i>	% in SEQ ID NO:5	% in <i>tcbA</i>	% in dicot	% in maize
Ala	GCT	62	21	69	41	42	24
	GCC	26	32	27	17	27	34
	GCA	11	25	4	22	25	18
	GCG	0	21	0	21	6	24
Arg	AGG	48	0	60	2	25	26
	CGC	22	36	18	16	11	24
	AGA	20	11	15	6	30	15
	CGT	11	39	7	57	21	11
	CGG	0	7	0	13	4	15
	CGA	0	8	0	6	8	9
Asn	AAC	100	32	100	33	55	68
	AAT	0	68	0	67	45	32
Asp	GAC	67	22	70	25	42	63

amino acid	codon	% in SEQ ID NO:3	% in tcdA	% in SEQ ID NO:5	% in tcbA	% in dicot	% in maize
	GAT	33	78	30	75	58	37
Cys	TGC	100	30	100	19	56	68
	TGT	0	70	0	81	44	32
End	TGA	100	0	100	0	33	59
	TAG	0	0	0	0	19	21
	TAA	0	100	0	100	48	20
Gln	CAA	65	61	74	53	59	38
	CAG	35	39	26	47	41	62
Glu	GAG	100	24	98	36	51	71
	GAA	0	76	2	64	49	29
Gly	GGT	67	37	64	44	33	20
	GGC	32	36	36	22	16	42
	·GGA	1	20	0	19	38	19
	GGG	0	8	0	16	12	20
His	CAC	62	40	72	31	46	62
	CAT	38	60	28	69	54	38
Ile	ATC	73	34	65	24	37	58
	ATT	27	51	35	59	45	28
	ATA	0	15	0	17	18	14
Leu	CTC	54	11	59	7	28	26
	TTG	29	17	25	32	26	15
	CTT	16	9	15	7	19	17
	TTA	0	18	0	19	10	5
	CTG	0	32	0	29	9	29
	CTA	0	13	0	7	8	8
Lys	AAG	99	79	99	75	61	78
	AAA	1	21	1	25	39	22
Met	ATG	100	100	100	100	100	100
Phe	TTC	100	42	100	41	55	71
	TTT	0	58	0	59	45	29
Pro	CCA	74	30	91	26	42	26
	CCT	22	28	7	20	32	22
	CCC	4	14	3	7	17	24
	CCG	0	27	0	47	9	28
Ser	TCC	47	19	55	11	18	23
	TCT	35	15	30	15	25	15
	AGC	18	22	15	18	18	23
	AGT	0	20	0	31	14	9
	TCG	0	7	0	8	6	14
	TCA	0	17	0	17	19	16
Thr	ACC	60	41	64	31	30	37
	ACT	28	25	32	34	35	20
	ACA	12	21	4	18	27	21
	ACG	0	13	0	18	8	22
Trp	TGG	100	100	100	100	100	100
Tyr	TAC	100	24	100	19	57	73
	TAT	0	76	0	81	43	27
Val	GTC	69	27	73	11	20	31
	GTG	21	17	22	27	29	39
	GTT	10	34	3	48	39	21
	GTA	0	22	2	14	12	8

EXAMPLE 1

Design Of Plant Codon-Biased Genes Encoding W-14 Peptides
TcbA and TcdA

A. Gene Design

The coding strands of the native DNA sequences of the *Photorhabdus* W-14 genes encoding peptides TcbA and TcdA were scanned for the presence of deleterious sequences such as the Shaw/Kamen RNA destabilizing motif ATTTA, 5 intron splice recognition sites, and poly A addition motifs. This was done using the MacVector Sequence Analysis Software (Oxford Molecular Biology Group, Symantec Corp.), using a custom Nucleic Acid Subsequence File. The native sequence was also searched for runs of 10 4 or more of the same base.

Motif searching of the native W-14 *tcbA* and *tcdA* genes revealed the presence of many potentially deleterious sequences in the protein coding strands, as summarized in Table 4. Not shown, but also present, were 15 many runs of four or more single residues (e.g. the native *tcbA* gene has 81 runs of four A's).

Table 4

Native Gene	ATTTA	5' Splice	3' Splice	Poly A Addition*	RNAP II term.
<i>tcbA</i>	18	7	17	46	0
<i>tcdA</i>	18	7	13	77	1

* Totals of 16 different motifs.

Analyses of eukaryotic genes and plant genes in 20 particular have shown that CG & TA doublets are underrepresented, while the genes are enriched in CT & TG doublets. The sequences of the hemicot biased genes have accordingly been adjusted to encompass these base 25 compositions and to have G+C compositions of about 53%, similar to many plant genes. When compared to the native W-14 *tcbA* and *tcdA* genes, the plant-biased genes have a much more uniform G+C distribution.

Nucleotide changes to remove potentially deleterious sequences were chosen to simultaneously adjust the codon 30 composition of the coding region to more closely reflect that of plant genes. A framework for these changes was provided by the codon bias tables prepared for maize and dicot genes shown in Table 3.

Comparison of codon compositions of the native W-14 genes to maize and dicot genes revealed that the W-14 genes contain a very different preference set of the degenerate codons for the 18 amino acids for which there is a choice (Table 3). For each of 8 amino acids (Phe, Tyr, Cys, Arg, Asn, Lys, Glu, and Gly) in both W-14 genes, the most abundant codon is different from the preferred codons found in either maize or dicot genes. One might expect that translational difficulties would be encountered in efforts to produce in plants proteins (such as *TcbA* and *TcdA*) having high relative amounts of these amino acids from mRNAs having large numbers of nonpreferred codons. There is a marked difference in distribution of the codon compositions specifying the other 10 amino acids. For His, Gln, Ile, Val, and Asp, the dicot-preferred codons are found as the most abundant ones in both W-14 genes. For Leu, Thr, Ser, and Ala, the maize preferred codons are the most abundant codon choices found in the *tcdA* gene. In contrast, the *tcbA* gene contains only the CCG (Pro) maize-preferred codon as the highest abundance choice.

In making the codon choices, doublet contents were considered, so that adjacent codons preferably did not form CG or TA doublets (which are underrepresented in eukaryotic genes; 1, 4), while CT or TG doublets (which are enriched in eukaryotic genes ibid.) were created when possible.

Choices were also made to utilize a diversity of codons for Met, Trp, Asn, Asp, Cys, Glu, His, Ile, Lys, Phe, Thr, and Tyr.

The sequences were also designed to encode unique 6-bp recognition sites for restriction enzymes, spaced about every 1200 bp. Finally, an additional codon (GCT; Ala) was inserted at the second position to encode an Nco I recognition site encompassing the ATG (Met) start codon. Additional recognition sites were included after

the stop codon to facilitate subsequent cloning steps into expression vectors. These features are set forth above in Tables 1 and 2.

The new *tcdA* and *tcbA* genes of SEQ ID NO:3 and SEQ 5 ID NO:4 share 73.5%, and 72.6% identity, respectively, to their native W-14 counterparts (Wisconsin Genetics Computer Group, GAP algorithm).

B. Gene Synthesis

The complete synthesis of the plant codon-biased 10 *tcbA* and *tcdA* genes was performed under contract by Operon Technologies, Inc. (OPTI, Alameda, CA). Basically, chemically synthesized oligonucleotides of appropriate sequence were assembled into DNA pieces about 500 bases long. These were joined together end-to-end 15 (presumably by means of appropriately placed restriction enzyme sites) into four larger pieces of roughly 2 kilobase pairs (kbp) each; therefore each comprised about 1/4 of the entire coding region of the particular gene. DNA sequence of the pieces was confirmed at this step. 20 If mistakes in sequence were present, the appropriate oligonucleotides were re-synthesized, and the assembly process was repeated. Once gene fractional parts were sequence verified, they were assembled in pairs to make the gene halves, and again sequence verified. Finally, 25 the two halves were joined, and the sequences of the junctions between the halves was verified. Therefore, each part of the new gene was sequence verified at least twice.

It should be noted that attempts to express the 30 native *tcbA* or *tcdA* genes in standard *Escherichia coli* cloning strains suggests that production of these proteins is lethal. Lethality problems may be encountered if standard cloning vectors having leaky expression from inherent *lacZ* promoters are used to 35 assemble these genes.

C. Addition Of Endoplasmic Reticulum Targeting Peptide To
Tcda Coding Region

It is known to those in the field of plant gene expression that proteins are specifically directed into the endoplasmic reticulum (ER) by means of a short signal peptide which is removed during or after the transport process through the ER membrane. The mature (processed) protein is incorporated into the ER endomembrane or is released into the ER lumen where the transported protein may be uniquely folded (aided by chaperonins), modified by glycosylation, accumulated in the vacuole, or additionally translocated (by secretion). These processes are reviewed by Gomord and Faye [V. Gomord and L. Faye, (1996) *Signals and mechanisms involved in intracellular transport of secreted proteins in plants*. Plant Physiol. Biochem. 34:165-181] and by Bar-Peled et al. [M. Bar-Peled, D. C. Bassham, and N. V. Raikhel, (1996) *Transport of proteins in eukaryotic cells: more questions ahead*. Plant Molec. Biology 32:223-249]. It is also known that the subcellular recognition mechanisms for an ER signal peptide are evolutionarily somewhat conserved, since the ER signal for a protein normally produced in monocot (maize) cells is recognized and processed normally by dicot (tobacco) cells. This is exemplified by the maize 15 kDa zein ER signal peptide [L. M. Hoffman, D. D. Donaldson, R. Bookland, K. Rashka, and E. M. Herman, (1987) *Synthesis and protein body deposition of maize 15-kd zein in transgenic tobacco seeds*. EMBO J. 6:3213-3221, and U.S. Patent 5589616]. Further, it is known that the ER signal peptide derived from one protein can direct the translocation of a different protein if it is appropriately attached to the second protein by genetic engineering methods [D. C. Hunt and M. J. Chrispeels, (1991) *The signal peptide of a vacuolar protein is necessary and sufficient for the efficient secretion of a cytosolic protein*. Plant

Physiol. 96:18-25, and Denecke, J., J. Botterman, and R. Deblaere (1990) *Protein secretion in plants can occur via a default pathway*. Plant Cell 2:51-59]. Therefore, one may expose a protein *in vivo* to different biochemical environments by directing its accumulation in the cytosol (by not providing a signal peptide sequence), or in the ER/vacuole (by provision of an appropriate signal peptide.)

The ER signal peptide of maize 15 kDa zein proteins is known to comprise the first 20 amino acids encoded by the zein coding region. Two examples of such signal peptides the ER signal peptide of 15 kDa zein from A5707 maize, NCBI Accession # M72708, and the ER signal peptide of 15 kDa zein from Black Mexican Sweet maize, NCBI Accession # M13507. There is only a single amino acid difference (Ser vs Cys at residue 17) between these signal peptides.

SEQ ID NO:5 is a modified sequence coding the ER signal peptide of 15 kDa zein from Black Mexican Sweet maize. The modifications embodied in this sequence were made to accommodate the different monocot/dicot codon usages and other sequence motif considerations discussed above in the design of the plant-optimized *tcdA* coding region. The sequence includes an additional Ala residue at position #2 to accommodate the *NcoI* site which spans the ATG start codon.

SEQ ID NO:6 gives a sequence coding for the full-length native *TcdA* protein (amino acids 22-2537) fused to the modified 15 kDa zein endoplasmic reticulum signal peptide (amino acids 1-21).

Example 2

Transformation Of Tobacco With *Agrobacterium* Carrying Plasmid pDAB2041 Encoding *Photorhabdus* Toxins

A. Plasmid pDAB2041

Preparation of tobacco transformation vectors was accomplished in three steps. First, a modified plant-optimized *tcdA* coding region was ligated into a tobacco

plant expression cassette plasmid. In this step, the coding region was placed under the transcriptional control of a promoter functional in tobacco plant cells. RNA transcription termination and polyadenylation were 5 mediated by a downstream copy of the terminator region from the *Agrobacterium* nopaline synthase gene. Two plasmids designed to function in this role are pDAB1507 and pDAB2006. In the second step, the complete gene comprised of the promoter, coding region, and terminator 10 region was ligated between the T-DNA borders of an *Agrobacterium* binary vector, pDAB1542. Also positioned between the T-DNA borders was a plant selectable marker gene to allow selection of transformed tobacco plant cells. In the third step, the engineered binary vector 15 plasmid was conjugated from its *E. coli* host strain into a disabled *Agrobacterium tumefaciens* strain capable of transforming tobacco plant cells that regenerate into fertile transgenic plants.

It is a feature of plasmid pDAB1507 that any coding 20 region having an *Nco*I site at its 5' end and a *Sac*I site 3' to the coding region, when cloned into the unique *Nco*I and *Sac*I sites of pDAB1507, is placed under the transcriptional control of an enhanced version of the CaMV 35S promoter. It is also a feature of pDAB1507 that 25 the 5' untranslated leader (UTR) sequence preceding the *Nco*I site comprises a modified version of the 5' UTR of the MSV coat protein gene, into which has been cloned an internally deleted version of the maize *Adh1S* intron 1. Additionally it is a feature of pDAB1507 that 30 transcription termination and polyadenylation of the mRNA containing the introduced coding region are mediated by termination/Poly A addition sequences derived from the nopaline synthase (Nos) gene. Finally, it is a feature of pDAB1507 that the entire assembly of promoter/coding 35 region/3'UTR can be obtained as a single DNA fragment by cleavage at the flanking *Not*I sites.

It is a feature of plasmid pDAB2006 that any coding region having an *Nco*I site at its 5' end and a *Sac*I site 3' to the coding region, when cloned into the unique *Nco*I and *Sac*I sites of pDAB2006, is placed under the transcriptional control of the CaMV 35S promoter. It is also a feature of pDAB2006 that the 5' untranslated leader (UTR) sequence preceding the *Nco*I site comprises a polylinker. Additionally it is a feature of pDAB2006 that transcription termination and polyadenylation of the mRNA containing the introduced coding region are mediated by termination/Poly A addition sequences derived from the nopaline synthase (Nos) gene. Finally, it is a feature of pDAB2006 that the entire assembly of promoter/coding region/3'UTR can be obtained as a single DNA fragment by cleavage at the flanking *Not*I sites.

It is a feature of pDAB1542 that any DNA fragment flanked by *Not*I sites can be cloned into the unique *Not*I site of pDAB1542, thus placing the introduced fragment between the T-DNA borders, and adjacent to the neomycin phosphotransferase II (kanamycin resistance) gene.

To prepare a plant-expressible gene to produce the non-targeted *TcdA* protein in tobacco plant cells, DNA of a plasmid (pAOH_4-OPTI) containing the plant-optimized *tcdA* coding region, (SEQ ID No:3) was cleaved with restriction enzymes *Nco*I and *Sac*I, and the large 7550 bp fragment was ligated to similarly-cut DNA of plasmid pDAB1507 to produce plasmid pDAB2040. DNA of pDAB2040 was then digested with *Not*I, and the 8884 bp fragment was ligated to *Not*I digested DNA of pDAB1542 to produce plasmid pDAB2041. This plasmid was then conjugated by triparental mating [Firoozabady, E., D. L. DeBoer, D. J. Merlo, E. L. Halk, L. N. Amerson, K. E. Rashka, and E. E. Murray (1987) *Transformation of cotton (Gossypium hirsutum L.) by Agrobacterium tumefaciens and regeneration of transgenic plants.* Plant Molec. Biol.

10:105-116] from the host *Escherichia coli* strain (XL1-Blue, Stratagene, La Jolla, CA), into the nontumorigenic *Agrobacterium tumefaciens* strain EHA101S, which is a spontaneous streptomycin-resistant mutant of strain 5 EHA101 (Hood, E. E., G. L. Helmer, R. T. Fraley, and M.-D. Chilton (1986) *The hypervirulence of Agrobacterium tumefaciens A281 is encoded in a region of pTiBo542 outside of T-DNA.* J. Bacteriol. 168:1291-1301). Strain EHA101S(pDAB2041) was then used to produce transgenic 10 tobacco plants that expressed the TcdA protein.

B. Plasmid pRK2013

To prepare a plant-expressible gene to produce the endoplasmic reticulum-targeted TcdA protein in tobacco plant cells, DNA of a plasmid (pAOH_4-ER) containing the 15 plant-optimized, ER-targeted *tcdA* coding region, (SEQ ID No:6) was cleaved with restriction enzymes *Nco*I and *Sac*I, and the large 7610 bp fragment was ligated to similarly-cut DNA of plasmid pDAB2006 to produce plasmid pDAB1833. DNA of pDAB1833 was then digested with *Not*I, and the 8822 20 bp fragment was ligated to *Not*I digested DNA of pDAB1542 to produce plasmid pDAB2052. This plasmid was then conjugated by triparental mating from the host *Escherichia coli* strain (XL1-Blue), into the nontumorigenic *Agrobacterium tumefaciens* strain EHA101S. 25 Strain EHA101S(pDAB2052) was then used to produce transgenic tobacco plants that expressed the TcdA protein containing an amino terminus endoplasmic reticulum targeting peptide.

30 C. Transfer of Plasmid pDAB2041 Into *Agrobacterium* Strain EHA101S

Cultures of *E. coli* carrying the engineered Ti plasmid pDAB2041 (plasmid containing the rebuilt Toxin A gene, *tcdA*), *E. coli* carrying the plasmid pRK2013, and 35 *Agrobacterium* strain EHA101S were grown overnight, then mixed 1:1:1 on plain LB medium solidified with agar and

cultured in the dark at 28°C. Two days later, the lawn of bacteria was scraped up with a loop, suspended in plain LB medium, vortexed, and then diluted 1:10⁴, 1:10⁵, and 1:10⁶ fold in plain LB liquid medium. Aliquots of these 5 dilutions were spread on selective plates containing medium YEP plus erythromycin (100 mg/L) and streptomycin (250 mg/L) and grown at 28°C. Two days later, single colonies were picked and streaked onto the same medium, then spread to give single colonies. Single colonies were 10 picked again and streaked, then spread for single colonies. Single colonies were picked a third time, grown as streaks, then subjected to a quality analysis involving growth on lactose medium and chromogenic assay with Benedict's reagent. Of ten strains developed in this 15 way, the fastest coloring colony was chosen for further work.

D. Transformation Of Tobacco With *Agrobacterium* Carrying Plasmid pDAB2041

20 Tobacco transformation with *Agrobacterium tumefaciens* was carried out by a method similar, but not identical, to published methods (R Horsch et al, 1988. Plant Molecular Biology Manual, S. Gelvin et al, eds., Kluwer Academic Publishers, Boston). To provide source 25 tissue for the transformation, tobacco seed (*Nicotiana tabacum* cv. Kentucky 160) were surface sterilized and planted on the surface of TOB-, which is a hormone-free Murashige and Skoog medium (T. Murashige and F. Skoog, 1962). A revised medium for rapid growth and bioassays 30 with tobacco tissue culture. Plant Physiol. 75: 473-497) solidified with agar. Plants were grown for 6-8 weeks in a lighted incubator room at 28-30°C and leaves were collected sterilely for use in the transformation protocol. Approximately one cm² pieces were sterilely cut 35 from these leaves, excluding the midrib. Cultures of the

Agrobacterium strains (EHA101S containing pDAB2041), which had been grown overnight on a rotor at 28°C, were pelleted in a centrifuge and resuspended in sterile Murashige & Skoog salts, adjusted to a final optical 5 density of 0.7 at 600 nm. Leaf pieces were dipped in this bacterial suspension for approximately 30 seconds, then blotted dry on sterile paper towels and placed right side up on medium TOB+ (Murashige and Skoog medium containing 1 mg/L indole acetic acid and 2.5 mg/L 10 benzyladenine) and incubated in the dark at 28°C. Two days later the leaf pieces were moved to medium TOB+ containing 250 mg/L cefotaxime (Agri-Bio, North Miami, Florida) and 100 mg/L kanamycin sulfate (AgriBio) and incubated at 28-30°C in the light. Leaf pieces were moved 15 to fresh TOB+ with cefotaxime and kanamycin twice per week for the first two weeks and once per week thereafter. Leaf pieces which showed regrowth of the Agrobacterium strain were moved to medium TOB+ with cefotaxime and kanamycin, plus 100 mg/l carbenicillin 20 (Sigma). Four to six weeks after the leaf pieces were treated with the bacteria, small plants arising from transformed foci were removed from this tissue preparation and planted into medium TOB- containing 250 mg/L cefotaxime and 100 mg/L kanamycin in Magenta GA7 25 boxes (Magenta Corp., Chicago). These plantlets were grown in a lighted incubator room. After 3-4 weeks the primary transgenic plants had rooted and grown to a size sufficient that leaf samples could be analyzed for expression of protein from the transgene. Twenty-five 30 independent transgenic events were recovered as single plants from the pDAB2041 transformation.

Eight independent lines expressing various levels of transgenic protein from the T-DNA of pDAB2041 were propagated *in vitro* from leaf pieces as follows. Twelve 35 to sixteen approximately one cm² pieces were steriley cut from leaves of each primary transgenic plant, excluding

the midrib and all naturally occurring edges. These leaf pieces were placed on medium TOB+ containing 250 mg/L cefotaxime and 100 mg/L kanamycin, and cultured in the lighted incubator at 28-30°C for 3-4 weeks, at which time 5 small plants could be cut from the proliferating tissue mass. Several small plantlets from each transgenic line were moved into Magenta boxes containing medium TOB- plus cefotaxime and kanamycin and allowed to root and grow. The proliferating tissue mass was further cultured on 10 medium TOB+ with cefotaxime and kanamycin, and additional plants could be cut out and grown up as needed.

Plants were moved into the greenhouse by washing the agar from the roots, transplanting into soil in 5 $\frac{1}{2}$ " square pots, placing the pot into a Ziploc bag 15 (DowBrands), placing plain water into the bottom of the bag, and placing in indirect light in a 30°C greenhouse for one week. After one week the bag could be opened; the plants were fertilized and allowed to grow further, until the plants were acclimated and the bag was removed. 20 Plants were grown under ordinary warm greenhouse conditions (30°C, 16 H light). Plants were suitable for sampling four weeks post transplant.

Example 3

25 Characterization Of Transgenic Tobacco Plants Expressing *Photorhabdus* Toxin That Confer Insect Control.

A. Polyclonal Antibody Production

The *E. coli* produced recombinant TcdA protein was 30 purified by a series of column purification. The protein was sent to Berkley Antibody Company (Richmond, CA) for the production of antiserum in a rabbit. Inoculations with the antigen were initiated with 0.5 mg of protein followed by four boosting injections of 0.25 mg each at 35 about three week intervals. The rabbit serum was tested by the standard Western analysis using the recombinant TcdA protein as the antigen and enhanced chemi-

luminescens, ECL method (Amersham, Arlington Heights, IL) . The antibodies (PAb-EA₀) were purified using a PURE I antibody purification kit (Sigma, St. Luis, MO). PAb-EA₀ antibodies recognize the full-length TcdA and its 5 processed components.

B. Expression Of TcdA Protein In Tobacco

Protein was extracted from the leaf tissue of transformed and non-transformed tobacco plants following the procedure described immediately below.

10 Two leaf disks of 1.4 cm in diameter were harvested from the middle portion of a fully expanded leaf. The disks were placed on a 1.6 x 4 cm piece of 3M Whatman paper. The paper was folded lengthwise and inserted in a flexible straw. Four hundred micro liters of the 15 extraction buffer (9.5 ml of 0.2 M NaH₂PO₄, 15.5 ml of 0.2 M Na₂HPO₄, 2 ml of 0.5 M Na₂EDTA, 100 ml of Triton X100, 1 ml of 10% Sarkosyl, 78 ml of beta-mercaptoethanol, H₂O to bring total volume to 100 ml) was pipetted on to the paper. The straw containing the sample was then passed 20 through a rolling device used for squeezing out the extract 1.5 mL micro centrifuge tube was placed at the other end of the straw to collect the extract. The extract was centrifuged for 10 minutes at 14,000 rpm in an Eppendorf regrigerated microcentrifuge. The 25 supernatant was transferred into a new tube. Protein quantitation analysis was performed using the standard Bio-Rad Protein Analysis protocol (Bio-Rad Laboratories, Hercules, CA). The extract was diluted to 2 mg/ml of total protein using the extraction buffer.

30 For the detection of transgenic protein, Western blot analysis was performed. Following a standard procedure for protein separation (Laemmli, 1970), 40 µg of protein was loaded in each well of 4-20% gradient polyacrylamide gel (Owl Scientific Co., MA) for 35 electrophoresis. Subsequently, the protein was

transferred onto a nitrocellulose membrane using a semi-dry electroblotter (Pharmacia LKB Biotechnology, Piscataway, NJ). The membrane was incubated for one hour in Blotto (5% milk in TBST solution; 25 mM Tris HCL pH 5.4, 136 mM NaCl, 2.7 mM KCl, 0.1% Tween 20). Thereafter, Blotto was replaced by the primary antibody solution (in Blotto). After one hour in the primary antibody, the membrane was washed with TBST for five minutes three times. Then the secondary antibody in Blotto (1:2000 dilution of goat anti-rabbit IgG conjugated to horseradish peroxidase; Bio-Rad Laboratories) was added to the membrane. After one hour of incubation, the membrane was washed with an excess amount of TBST for 10 minutes four times. The protein was visualized by using the enhanced chemi-luminescens, ECL method (Amersham, Arlington Heights, IL). The differential intensity of the protein bands were measured using densitometer (Molecular Dynamics Inc., Sunnyvale, CA).

To determine the expression of TcdA protein in tobacco transformed with pDAB2041, PAb-EA₀ antibodies were used as the primary antibodies. The expression levels of TcdA protein varied among independent transformation events. The primary plant generated from the event #2041-13 showed the highest level of pre-pro TcdA expression of extractable protein. When the leaf pieces from this plant (#2041-13) were used in *in vitro* propagation, several plants were obtained. Seven of these plants were analyzed for the expression of the TcdA protein. All but one plant produced the full-length TcdA protein as well as some processed peptide components. Using the antibodies specific to Neomycin phosphotransferase, NPT (5 prime-3 prime, Boulder, Co), the expression the selectable marker gene (*npt II*) was detected. Similar results were obtained for #2041-29.

Western analysis of plants derived from event #2041-13.

Plant #	TcdA	NPT (selectable marker)
2041-13A	+	not done
2041-13B	+	not done
2041-13-1	-	+
2041-13-2	+	+
2041-13-3	+	+
2041-13-4	+	+
2041-13-5	+	+

C. Nucleic Acid Analysis of Transgenic Tobacco Lines

Genomic DNA was prepared from a group of 2041 5 transgenic events. The lines included Magenta box stage 2041-13, and greenhouse stage plants 2041-13-1, 2041-13-2, 2041-13-5, 2041-9, 2041-20A and 2041-20B. A transgenic GUS line (2023) was included as a negative control. Southern analysis of these lines was performed. 10 The genomic tobacco DNA was restricted with the enzyme SstI which should result in a 8.9 kb hybridization product when hybridized to a *tcdA* gene specific probe. The 8.9 kb hybridization product should consist of the 35T promoter and the *tcdA* coding region. All 2041 plants 15 contained a band of the expected size. Events 2041-9 and -20 appear to be the same line with 5 identical hybridizing bands. Event 2041-13 produced 6 hybridization fragments with the *tcdA* coding region probe. Magenta box and various greenhouse plants of 20 2041-13 all produced the same hybridization profile. This hybridization pattern was different from that of events 2041-9 and -20.

RNA analysis, using the *tcdA* coding region probe, was performed on the same group of greenhouse 2041 25 plants. Immunoblot analysis had revealed that plants 2041-9, 2041-20A, 2041-20B, and 2041-13-1 produced no detectable TcdA protein; while 2041-13-2 and 2041-13-5 produced substantial amounts of full-length TcdA. Northern analysis was in agreement with the immunoblot.

result. A faint RNA signal was detected for plants 2041-9, 2041-20A, 2041-20B, and 2041-13-1. Only faintly visible was a band corresponding to full-length *tcdA* transcript in plant 2041-13-1. In contrast, for plants 5 2041-13-2 and 2041-13-5 a strong RNA signal was detected, with a substantial amount of full-length size (~8.0 kb) *tcdA* transcript. These data support the observed bioassay activity for this group of plants.

Genomic DNA was prepared from a second functionally 10 active 2041 transgenic event, 2041-29. Southern analysis of this line was performed. A transgenic GUS line (2023) was included as a negative control, DNA of line 2041-9 was included as a positive control.

The genomic tobacco DNAs were restricted with the 15 enzyme *Sst*I which should result in a 8.9 kb hybridization product when hybridized to a *tcdA* gene specific probe. The 8.9 kb hybridization product should consist of the 35T promoter and the *tcdA* coding region. For plant 2041-29-5, three hybridization products larger than 8.9 kb the 20 were detected with the *tcdA* gene specific probe.

Immunoblot analysis has demonstrated pre-pro TcdA protein is made by this plant, it is therefore likely that a restriction site was lost during transformation or regeneration, or the 2041-29 genomic DNA was not 25 thoroughly digested.

D. Tobacco Leaf-Disk Tests With Tobacco Hornworm Exhibiting Insect Control

Leaves were sampled from tobacco plants, *Nicotiana* 30 *tabaco*, previously transplanted into the greenhouse. A single leaf was sampled from each plant on each test date. Leaves were selected from the zone where younger elongate leaves transition into older ovate leaves. Excised leaves were placed into 12 oz. cups with the 35 petiole submerged in water to maintain turgor, and transported to the laboratory.

Eight, 1.4 cm disks were cut from the center portion of one side of each leaf (right adaxial side up, with distal portion facing away from the observer). Each disk was placed individually into a well of a C-D

5 International 128 well tray (Pitman, NJ.) into which 0.5 ml of a 1.6% aqueous agar solution had been previously pipetted. The solidified agar prevented the leaf disks from drying out. The adaxial surface of the disk was always oriented up.

10 A single neonate tobacco hornworm, *Manduca sexta*, was placed on each disk and the wells were sealed with vented plastic lids. The assay was held at 27°C and 40% RH. Larval mortality and live-weight data were collected after 3 days. Data were subjected to analysis of

15 variance and Duncan's multiple range test ($\alpha = 0.05$) (Proc GLM, SAS Institute Inc., Cary, NC.). Data were transformed using a logarithmic function to correct a correlation between the magnitude of the mean and variance.

20

Table 6
Results of leaf-disk assays from greenhouse grown tobacco plants with event 2041-13.

TRT	Plant	Plant Age	Weight of Surviving Larvae (mg) & Duncan's Group ¹				
			Pretest	Test 1	Test 2	Test 3	3 Test Sum.
13	non-transformed - 2	young	---	---	---	18.8 a*	---
14	non-transformed - 3	young	---	---	---	17.0 ab	---
16	non-transformed - 5	young	---	---	---	16.4 ab	---
3	2041-13-1 (western -)	young	---	17.6 a	18.2 a	16.1 ab	17.3 a
9	Gus Control	old	19.3 a	14.6 a	16.3 a	14.5 ab	15.1 a
10	non-transformed - 1	young	---	8.3 b	16.8 a	13.9 b	13.0 b
11	2041-20B (western -)	old	---	10.0 b*	13.7 ab	14.6 ab	12.9 b
15	non-transformed - 4	young	---	---	---	13.0 bc	---
8	2041-20A (western -)	old	15.7 a	8.3 b	11.3 bc	9.2 cd	9.6 c
12	2041-9 (western -)	old	19.5 a	---	---	7.9 d	---
7	2041-13-5 (western +)	young	---	6.3 bc	9.6 cd	7.2 de	7.7 d
5	2041-13-3 (western +)	young	---	6.4	6.2 e	6.8 de**	6.4 de
				bc****			
1	2041-13A (western +)	old	7.2 b	6.8 bc*	7.0 de*	5.4 e	6.4 de
6	2041-13-4 (western +)	young	---	4.9 c****	5.8 e	7.6 d	6.4 de
4	2041-13-2 (western +)	young	---	5.7 bc	5.7 e**	7.5 d	6.3 de
2	2041-13B (western +)	old	---	4.7 c**	5.6 e	7.2 de	5.9 e

* Number of stars corresponds to the number of dead larvae per 8 tested.

1. Data transformed (logarithm) for analysis.

Means followed by the same letter are not significantly different (alpha = 0.05).

5

TABLE 7
Results Of Leaf-Disk Assays From Greenhouse Grown Tobacco Plants
With Event 2041-29.

Plant	MEAN WGT (MG) / Duncan's Group				
	Test 1	Test 2	Test 3	Test 4	Four Test Summary
2014-6 GUS 1	15.8 a	16.6 a	**5.5bc	*12.9ab	13.2 a
2014-6 GUS 2	14.4 a	*6.6 bc	*13.4a	15.2a	12.6 a
KY-160 NTC	13.4 a	6.7 bc	7.9b	8.5bc	9.1 b
2041-29 4P	*4.9 b	*7.3b	****6.9b	*****	6.3 c
2041-29 7	*5.9 b	5.1bc	***6.7b	***7.2c	6.1 c
2041-29 3P	*5.6 b	**7.9b	*****6.5b	***3.6d	5.9 c
2041-29 2P	6.3 b	****4.7c	*****4.1c	*****4.6d	5.4 c

* Number of stars corresponds to the number of dead 10 larvae per 8 tested.

1. Data transformed (logarithm) for analysis.

Means followed by the same letter are not significantly different (alpha = 0.05).

All event 2041-29 plants significantly depressed THW 15 larval weight gain compared to control plants. Average weight depression was 49%. Statistically significant mortality occurred in THW larvae exposed to foliage from 2041-29 plants. Mortality averaged 37.5% compared to 5.2% in controls.

20

E. Isolation and Characterization of Functional Photorhabdus Toxin Protein From Transgenic Plants

Seven grams of transgenic tobacco plants (2041-13) expressing TcdA (Toxin A) gene were homogenized with 10 25 ml 50 mM Potassium Phosphate buffer, pH 7.0 using a bead beater (Biospec Products, Bartlesville, OK) according to manufacturer's instructions. The homogenate was filtered through four layers of cheese cloth and then centrifuged at 35,000 g for 15 min. The supernant was collected and 30 filtered through 0.22 μ m Millipore ExpressTM membrane. It was then applied to a Superdex 200 column (2.6 x 40 cm)

which had been equilibrated with 20 mM Tris buffer, pH 8.0 (Buffer A). The protein was eluted in Buffer A at a flow rate of 3 ml/min. Fractions with 3 ml each were collected and subjected to southern corn rootworm (SCR) 5 bioassay. It was found that fractions corresponding to a native molecular weight around 860 kDa had the highest insecticidal activity. Western analysis of the active fraction using a polyclonal antibody specific to Toxin A indicated the presence of full-length TcdA peptide. The 10 active fractions were further combined and applied to a Mono Q 10/10 column which had been equilibrated with Buffer A. Proteins bound to the column were then eluted by a linear gradient of 0 to 1 M NaCl in Buffer A. Fractions with 2 ml each were collected and analyzed by 15 both SCR bioassay and Western using antibody specific to Toxin A. The results again demonstrated the correlation between insecticidal activity and presence of full-length TcdA peptide.

20 F. Characterization of Progeny Transgenic Plants

The inheritability of the genetically engineering plants containing the *Photobrhabdus* toxin gene was evaluated by generating F1 progeny. Progeny was generated from 2041-13 event by selfing expression 25 positive plants. The 2041-13 plants in the greenhouse were allowed to self-pollinate. Seed capsules were collected when mature and were allowed to dry and after-ripen on the laboratory bench for two weeks. Seed from plant designated 2041-13A was surface-sterilized and 30 distributed on the surface of medium TOB- without selection, to allow recovery of nonexpressing or nontransgenic progeny as well as expressing and segregating transgenic siblings. Seed was germinated in a C lighted incubator room (16 H light, 28 C). After 1 35 month, fifty-one seedlings, designated 2041-13A-S1 through S51, were distributed into Magenta boxes

containing medium TOB- to grow further. Three weeks later, leaf samples from these Magenta-box grown seedlings were submitted for evaluation of the level of expression of TcdA toxin.

5 Leaf samples were tested for kanamycin response by placing sterile leaf segments on medium TOB+ containing 100 mg/L kanamycin in the light and scoring for tissue growth and color after two weeks. All leaf pieces showed some positive response, indicating complex segregation.

10 This group of in vitro grown event 2041-13 progeny seedlings were all transplanted into the greenhouse approximately two months after seeding onto medium, using the following method. After washing the agar from the roots, plants were transplanted into 5 ½ inch square pots
15 in a soil mix containing 75% MetroMix and 25% mineral soil. They were enclosed in a zip-lock bag and plain water added to leave 1-2 inches of water in the bottom of the bag after soil absorption. These bags were closed and placed under a cart in the greenhouse to protect them
20 from direct sunlight. The bags were opened after 5-6 days, and removed after 7 days, when the plants were adapted to soil and were moved to the top of the cart for normal greenhouse culture. Plants were ready to test in insect bioassays at four weeks post transplant.

25 F1 progeny were evaluated for expression of protein toxin by immunological screen and for biological activity by plant bioassays, as described previously, using tobacco hornworm. There existed a positive correlation between levels of expression protein toxin and degree of
30 growth inhibition and at higher expression levels mortality was observed. The biological activity was observed to be statistical significance with high confidence levels between populations of non-transformed and transformed expressing protein toxin.

35 The following table summarizes the results of insect (tobacco hornworm) bioassays conducted with F1 progeny of

self-fertilized 2041-13 plants genetically engineered to produce the "204" A toxin. The tests included 6 non-expressing progeny (protein-negative controls), 45 toxin A expressors, and 4 non-transformed controls (KY-160).

5 Results are from three leaf-disk assays (method previously outlined) where eight disks were used per test. The data were analyzed using analysis of variance and were blocked by test.

The treatment effect for each of these analyses 10 indicated the $Pr > F$ was less than 0.0001. The Toxin A expressors produced significant control of tobacco hornworm compared to each of the control groups based on each of the three measures of efficacy. The two control groups behaved similarly. Statistical analysis using 15 ANOVA and an LSD test with alpha equal to 0.01 (or 1%) showed differences between the 3 groups. The LSD test indicated that the non-expressors and the non-transformed plants were similar in larvae weights but the expressors gave weights significantly lower than either of the other 20 two groups of plants. These data demonstrated that the genetic basis for insect control was inheritable and corresponded to the presence of expressed toxin gene.

Table 8
Tobacco hornworm results from F1 progeny of self-fertilized 2041-13 tobacco plants.

Treatment Group	Mean Value and Duncan's Grouping ^d		
	Total Weight (mg) ^a	Survivor Weight (mg) ^b	Leaf Area (cm ²) ^c
Non-transformed Control	15.8 a	15.8 a	1.2 a
Protein-negative Control	16.4 a	16.5 a	1.2 a
Toxin A Expressor	8.1 b	9.2 b	4.9 b

^a Average insect weight with dead insects considered to weigh nothing.

^b Average insect weight with dead insects excluded from 30 analysis.

^c Total leaf area remaining per eight leaf disks. Initial area was approximately 12 cm².

^d Means followed by the same letter are not significantly different (alpha = 0.05).

Example 4

5 Transformation Of Maize With a Vector Carrying Plasmid
pDAB1834 Encoding *Photorhabdus* ToxinsA. Preparation Of Maize Transformation Vectors
Containing Modified Plant-Optimized *Tcda* Coding Regions:
Plasmid Pdab1834

10

Preparation of maize transformation vectors was accomplished in two steps. First, a modified plant-optimized *tcda* coding region was ligated into a plant expression cassette plasmid. In this step, the coding 15 region was placed under the transcriptional control of a promoter functional in maize plant cells. RNA transcription termination and polyadenylation were mediated by a downstream copy of the terminator region from the *Agrobacterium* nopaline synthase gene. One 20 plasmid designed to function in this role is pDAB1538. In the second step, the complete gene comprised of the promoter, coding region, and 3' UTR terminator region was ligated to a plant transformation vector that contained a plant expressible selectable marker gene which allowed 25 the selection of transformed maize plant cells amongst a background of nontransformed cells. An example of such a vector is pDAB367.

It is a feature of plasmid pDAB1538 that any coding 30 region having an *Nco*I site at its 5' end and a *Sac*I site 3' to the coding region, when cloned into the unique *Nco*I and *Sac*I sites of pDAB1538, is placed under the transcriptional control of the maize ubiquitin1 (ubil) promoter. It is also a feature of pDAB1538 that the 5' 35 untranslated leader (UTR) sequence preceding the *Nco*I site comprises a polylinker. Additionally it is a feature of pDAB1538 that transcription termination and polyadenylation of the mRNA containing the introduced coding region are mediated by termination/Poly A addition

sequences derived from the nopaline synthase (Nos) gene. Finally, it is a feature of pDAB1538 that the entire assembly of promoter/coding region/3'UTR can be obtained as a single DNA fragment by cleavage at the flanking *NotI* sites.

It is a feature of pDAB367 that the phosphinothricin acetyl transferase protein, which has as its substrate phosphinothricin and related compounds, is produced in plant cells through transcription of its coding region mediated by the Cauliflower Mosaic Virus 35S promoter and that termination of transcription plus polyadenylation are mediated by the nopaline synthase terminator region. It is further a feature of pDAB367 that any DNA fragment containing flanking *NotI* sites can be cloned into the unique *NotI* site of pDAB367, thus physically linking the introduced DNA fragment to the aforementioned selectable marker gene.

To prepare a maize plant-expressible gene to produce the endoplasmic reticulum-targeted TcdA protein in plant cells, DNA of a plasmid (pAOH_4-ER) containing the plant-optimized, ER-targeted *tcdA* coding region, (SEQ ID No:6) was cleaved with restriction enzymes *NcoI* and *SacI*, and the large 7610 bp fragment was ligated to similarly-cut DNA of plasmid pDAB1538 to produce plasmid pDAB1832. DNA of pDAB1832 was then digested with *NotI*, and the 9984 bp *NotI* fragment was ligated into the unique *NotI* site of pDAB367 to produce plasmid pDAB1834.

It is a feature of plasmids pDAB1834 that the *ubil* and 35S promoters are encoded on the same DNA strand.

30

B. Transformation and Regeneration of Transgenic Maize Isolates

Type II callus cultures were initiated from immature zygotic embryos of the genotype "Hi-II." (Armstrong et al, (1991) Maize Genet. Coop. Newslett., 65: 92-93). Embryos were isolated from greenhouse-grown ears from

crosses between Hi-II parent A and Hi-II parent B or F_2 embryos derived from a self- or sib-pollination of a Hi-II plant. Immature embryos (1.5 to 3.5 mm) were cultured on initiation medium consisting of N6 salts and vitamins

5 (Chu et al, (1978) *The N6 medium and its application to anther culture of cereal crops*. Proc. Symp. Plant Tissue Culture, Peking Press, 43-56), 1.0 mg/L 2,4-D, 25mM L-proline, 100 mg/L casein hydrolysate, 10 mg/L $AgNO_3$, 2.5 g/L GELRITE (Schweizerhall, South Plainfield, NJ), and 20

10 g/L sucrose, with a pH of 5.8. After four to six weeks callus was subcultured onto maintenance medium (initiation medium in which $AgNO_3$ was omitted and L-proline was reduced to 6 mM). Selection for Type II callus took place for ca. 12-16 weeks.

15 Plasmid pDAB1834 was transformed into embryogenic callus. For blasting, 140 μ g of plasmid DNA was precipitated onto 60 mg of alcohol-rinsed, spherical gold particles (1.5 - 3.0 μ m diameter, Aldrich Chemical Co., Inc., Milwaukee, WI) by adding 74 μ L of 2.5M $CaCl_2$ H_2O and

20 30 μ L of 0.1M spermidine (free base) to 300 μ L of plasmid DNA and H_2O . The solution was immediately vortexed and the DNA-coated gold particles were allowed to settle. The resulting clear supernatant was removed and the gold particles were resuspended in 1 ml of absolute ethanol.

25 This suspension was diluted with absolute ethanol to obtain 15 mg DNA-coated gold/mL.

Approximately 600 mg of embryogenic callus tissue was spread over the surface of Type II callus maintenance medium as described herein lacking casein hydrolysate and L-proline, but supplemented with 0.2 M sorbitol and 0.2 M mannitol as an osmoticum. Following a 4 h pre-treatment, tissue was transferred to culture dishes containing blasting medium (osmotic media solidified with 20 g/L TC agar (PhytoTechnology Laboratories, LLC, Shawnee Mission, KS) instead of 7 g/L GELRITE. Helium blasting accelerated suspended DNA-coated gold particles towards

and into the prepared tissue targets. The device used was an earlier prototype of that described in US Patent 5,141,131 which is incorporated herein by reference.

5 Tissues were covered with a stainless steel screen (104 μm openings) and placed under a partial vacuum of 25 inches of Hg in the device chamber. The DNA-coated gold particles were further diluted 1:1 with absolute ethanol prior to blasting and were accelerated at the callus targets four times using a helium pressure of 1500 psi,

10 with each blast delivering 20 μL of the DNA/gold suspension. Immediately post-blasting, the tissue was transferred to osmotic media for a 16-24 h recovery period. Afterwards, the tissue was divided into small pieces and transferred to selection medium (maintenance

15 medium lacking casein hydrolysate and L-proline but containing 30 mg/L BASTA® (AgrEvo, Berlin, Germany)). Every four weeks for 3 months, tissue pieces were non-selectively transferred to fresh selection medium. After 7 weeks and up to 22 weeks, callus sectors found

20 proliferating against a background of growth-inhibited tissue were removed and isolated. The resulting BASTA®-resistant tissue was subcultured biweekly onto fresh selection medium. Following western analysis, positive transgenic lines were identified and transferred to

25 regeneration media. Western-negative lines underwent subsequent RNA spot blot analysis to identify negative controls for regeneration.

Regeneration was initiated by transferring callus tissue to cytokinin-based induction medium, which

30 consisted of Murashige and Skoog salts, hereinafter MS salts, and vitamins (Murashige and Skoog, (1962) *Physiol. Plant.* 15: 473-497) 30 g/L sucrose, 100 mg/L *myo*-inositol, 30 g/L mannitol, 5 mg/L 6-benzylaminopurine, hereinafter BAP, 0.025 mg/L 2,4-D, 30 mg/L BASTA®, and

35 2.5 g/L GELRITE at pH 5.7. The cultures were placed in low light (125 ft-candles) for one week followed by one

week in high light (325 ft-candles). Following a two week induction period, tissue was non-selectively transferred to hormone-free regeneration medium, which was identical to the induction medium except that it

5 lacked 2,4-D and BAP, and was kept in high light. Small (1.5-3 cm) plantlets were removed and placed in 150x25 mm culture tubes containing SH medium (SH salts and vitamins (Schenk and Hildebrandt, (1972) Can. J. Bot. 50:199-204), 10 g/L sucrose, 100 mg/L myo-inositol, 5 mL/L FeEDTA, and

10 2.5 g/L GELRITE, pH 5.8). Plantlets were transferred to 12 cm pots containing approximately 0.25 kg of METRO-MIX 360 (The Scotts Co. Marysville, OH) in the greenhouse as soon as they exhibited growth and developed a sufficient root system. They were grown with a 16 h photoperiod

15 supplemented by a combination of high pressure sodium and metal halide lamps, and were watered as needed with a combination of three independent Peters Excel fertilizer formulations (Grace-Sierra Horticultural Products Company, Milpitas, CA). At the 6-8 leaf stage, plants

20 were transplanted to five gallon pots containing approximately 4 kg METRO-MIX 360, and grown to maturity.

EXAMPLE 5

Characterization Of Transgenic Maize Plants

25 Expressing *Photorhabdus* Toxin That Confer Insect Control.

A. Insect Bioassays

A single leaf was sampled from each plant in each test. Eight, 1.4 cm disks were cut from the outer portion of each leaf (approximately 30cm long) avoiding the

30 center vein. Each disk was placed individually into a well of a C-D International 128 well tray (Pitman, NJ.) into which 0.5 ml of a 1.6% aqueous agar solution had been previously pipetted. The solidified agar prevented the leaf disks from drying out. The adaxial surface of

35 the disk was always oriented up.

Five neonate southern corn rootworms, *Diabrotica undecimpunctata howardi*, were placed on each disk and the wells were sealed with vented plastic lids. The assay was held at 27°C and 40% RH. Larval mortality and live-weight data were collected after 3 days. Data were subjected to analysis of variance and Duncan's multiple range test ($\alpha = 0.05$) (Proc GLM, SAS Institute Inc., Cary, NC.). Weight data were transformed using a logarithmic function to correct a correlation between the magnitude of the mean and variance.

TABLE 9
Results of Maize Leaf-disk Test vs SCR

Treatment	Mean % Kill (Duncan's)	Mean Survival Weight (mg) (Duncan's)
1834 - 11	68 A	0.064 A
1834 - 17	44 B	0.098 B
1834 - 15	26 BC	0.127 C
HiII control	13 C	0.161 C

Note: Means followed by the same letter are not significantly different based on Duncan's multiple range test ($\alpha=0.05$). Insect groups weighing less than 0.1 mg were set to 0.03 mg instead of zero to conduct a more conservative analysis. Mortality (arcsin(sqrt.)) and weight(log10) data were transformed for analyses.

The results shown in Table 9 demonstrated that two events expressing TcdA protein were statistically distinct from control lines bioassayed using SCR neonates by mortality and survival weight criteria. These results demonstrated that southern corn rootworm were functionally effected by feeding on maize plants containing and expressing the *tcdA* gene. Those plants from 1834-11 were used to generate progeny for testing of inheritability of transgene.

B. PRODUCTION AND PROGENY TEST OF *tcdA* TRANSGENIC MAIZE

Origin and growth of progeny plants: Sibling plants 1834-11-07 and 1834-11-08, clonally derived by regeneration 5 from the callus of transgenic maize event 1834-11, were transplanted to the greenhouse and pollinated with inbred OQ414. Seeds obtained from these crosses, comprising seed lots 1834-11-07A and 1834-11-08A, were planted in Roottrainers (1 ½ inch x 2 inch x 8 inch deep, product 10 #647, C. Hummert Intl., Earth City, Mo.) filled with Metro-Mix 360 soilless mix (Scotts Terra-Lite, available from Hummert Intl.) and top irrigated with Hoagland's nutrient solution. (Hoagland's solution contains 229 ppm nitrogen as nitrate, 24.6 ppm nitrogen as ammonium, 26 15 ppm P, 157 ppm K, 187 ppm Ca, 49 ppm Mg. and 30 ppm Na.)

Greenhouse conditions for this trial were: 16 hour days, daylight supplemented by metal halide lamps as needed to achieve a minimum of 600 ?Einstiens/cm² PAR, and ambient temperature 30 C days, 22 C nights.

20

Leaves were sampled for protein determination approximately one week after planting. Leaf bioassays were conducted 2-3 weeks after planting; root bioassays were initiated approximately 3 weeks post planting.

25

Protein analysis of progeny plants: Protein was extracted from leaf and root samples harvested from transgenic plants, line 1834-11 progenies, and non-transformed plants. Each sample was placed on a 1.6 x 4 cm piece of 30 3M Whatman™ paper. The paper was folded lengthwise and inserted in a flexible straw. A volume of 350 µl of an extraction buffer (9.5 ml of 0.2 M NaH₂PO₄, 15.5 ml of 0.2 M Na₂HPO₄, 2 ml of 0.5 M Na₂EDTA, 100 ml of Triton X-100, 1 ml of 10% Sarkosyl, 78 ml of beta-mercaptoethanol, H₂O 35 to bring total volume to 100 ml, 50 µg/ml Antipain, 50 µg/ml Leupeptin, 0.1 mM Chymostatin, 5 µg/ml Pepstatin) was pipetted on to the paper. The straw containing the

sample was then passed through a rolling device used for squeezing the extract into a 1.5 ml microcentrifuge tube. The extract was centrifuged for 10 minutes at 14,000 rpm in an Eppendorf refrigerated micro-centrifuge. The 5 supernatant was transferred into a new tube. The amount of the total extractable protein was determined using a standard BioRad Protein Analysis protocol (BioRad Laboratories, Hercules, CA).

The presence of the TcdA protein was visualized by 10 Western blot analysis following a standard procedure for protein separation (Laemmli, 1970). A volume of twenty μ l of extract was loaded in each well of 4-20% gradient polyacrylamide gel (Owl Scientific Co., MA) for electrophoresis. Subsequently, the protein was 15 transferred onto a nitrocellulose membrane using a semi-dry electroblotter (Pharmacia LKB Biotechnology, Piscataway, NJ). The membrane was incubated for one hour in TBST-M solution (10% milk in TBST solution; 25 mM Tris HCl pH 7.4, 136 mM NaCl, 2.7 mM KCl, 0.1% Tween 20). 20 Thereafter, the primary antibody (Anti-TcdA in TBST-M) was added. After one hour, the membrane was washed with TBST for five minutes, three times. Then the secondary antibody solution (goat anti-rabbit IgG conjugated to horseradish peroxidase; Bio-Rad Laboratories, in TBST-M) 25 was added to the membrane. After one hour of incubation, the membrane was washed with an excess amount of TBST for 10 minutes, four times. The protein was visualized using the Super Signal[®] West Pico chemiluminescence method (Pierce Chemical Co., Rockford, IL). The protein blot 30 was exposed on a Hyper-film (Amersham, Arlington Heights, IL) and was developed within 3 minutes. The intensity of the protein band was measured using a densitometer (Molecular Dynamics Inc., Sunnyvale, CA) and compared to standards.

35 Three of six plants from seed lot 1834-11-07A and three of six plants from seed lot 1834-11-08A produced

detectable levels of TcdA protein (Table 1).
Approximately 3.8 to 13.3 ppm of TcdA were detected in the leaf blades and 4.1 to 8.4 ppm were detected in the leaf tips of the protein-positive plants. The amounts of 5 TcdA protein detected in the roots were slightly lower than those found in the leaves.

Insect bioassays with progeny plants: Plants were selected for bioassay based on results from Western blot 10 analysis. Twelve (12), 6.4 mm diameter leaf discs were cut from the youngest leaf of each 2 week old seedling. Each disc was placed in a well of a 128-well tray (CD International) containing approximately 0.5mL of a solidified 2% agar in water solution. Two neonate 15 southern corn rootworm, *Diabrotica undecimpunctata howardi* (Barber) (SCR), were placed in each well with a leaf disc. Trays were covered with perforated lids and maintained under a controlled environment for 3 days (28 C; 16 hours light:8 hours dark; approx. 60% relative 20 humidity). Living larvae from 4 leaf discs were pooled and weighed producing 3 weight determinations per plant. Average weights were calculated by dividing the pooled 25 weight by the number of survivors. Differences in average weights of SCR fed leaf discs from protein positive and protein negative plants were assessed using analysis of variance on the natural log-transformed average weights (Minitab, v. 12.2, Minitab Inc., State College, PA).

30 Root bioassays were initiated approximately 1 week after the initiation of the leaf disc bioassays. Approximately 24h prior to eclosion, SCR eggs were suspended in a 0.15% solution of agar in water to a concentration of 100 eggs/ml. Plants were inoculated 35 with SCR eggs by pipetting 2.0 ml of the egg suspension (ie., approximately 200 eggs) just below the soil surface at the base of each plant. Two weeks after inoculation, plants were removed from their Rootrainer pots, their

roots washed free of potting mix, and scored for rootworm damage based on a 1 (resistant) to 9 (susceptible) rating system (Welch, 1977). The results of the root ratings were examined using non-parametric tests to determine if 5 the distribution of root ratings from the protein positive plants was the same as the distribution of the ratings from the protein negative plants. Testing was done at the 5% significance level. (StatXact v.3, CYTEL Software Corporation, Cambridge MA)

10

Results from leaf and root bioassays of tcdA protein positive and protein negative progeny plants are summarized in Table 10. The average weights of SCR larvae fed leaf discs from protein positive plants were 15 significantly lower than those of larvae fed leaf discs from protein negative plants ($F = 4.6$; $d.f. = 1, 34$; $P \leq 0.001$). The Kolmogorov-Smirnov 2 sample test ($p=0.04$) and the Wald Wolfowitz runs test ($p=0.001$) indicated that the protein positive and protein negative root rating 20 distributions were not similar. The Wilcoxon- Mann-Whitney test ($p=0.0206$) and the Normal Scores test ($p=0.206$) indicated that the average score for the protein positive plants was lower than the average root rating from the protein negative plants.

25

Table 10. Protein analysis and insect bioassay results with progeny of TcdA transgenic maize.

Plant Number	TcdA Protein	Leaf Disc Bioassay Avg. Wt. (mg)	Root Bioassay Root Rating (1-9)
1834-11-07A-30	PRO-	0.190	8
1834-11-08A-21	PRO-	0.196	9
1834-11-08A-16	PRO-	0.195	9
1834-11-08A-14	PRO-	0.137	9
1834-11-07A-22	PRO-	0.208	9
1834-11-07A-20	PRO-	0.175	9

1834-11-07A-26	PRO+	0.118	9
1834-11-08A-17	PRO+	0.132	8
1834-11-07A-14	PRO+	0.110	2
1834-11-07A-11	PRO+	0.106	4
1834-11-08A-28	PRO+	0.129	8
1834-11-08A-27	PRO+	0.108	4

DNA analysis of progeny plants: Leaf samples from 1834-11.7A and 1834-11.8A progeny plants were in conical 50 ml polypropylene tubes and dried in a Labconco Freeze Dry

5 Lyophilizer (Kansas City, MO) for 1-2 days. Lyophilized leaves were then ground in a Tecator Cyclotec 1093 Sample mill grinder (Hoganas, Sweden) and stored at -20C. Genomic DNA was extracted by the following procedure: (1) to a 25 ml Conical tube containing 300-500 mg of ground

10 tissue, 9 ml of CTAB (cetyl trimethylammonium bromide solution) was added, and incubated at 65°C for 1 hour; (2) 4.5 ml of chloroform: octanol (24:1) was added and mixed gently for 5 minutes; (3) samples were centrifuged at 2000 rpm and DNA was precipitated from the supernatant

15 with an equal volume of isopropanol; (4) DNA was collected on a glass hook, washed in ethanol, and dissolved in TE (10 mM Tris.HCl, 0.5 mM EDTA, pH8.0).

Genomic DNA was digested at 37 °C. for 2 hours in an

20 Eppendorf tube containing the following mixture:

8 µl of 800ug/ml DNA, 2 µl 1 mg/ml BSA (Bovine serum albumin), 2 µl 10x buffer, 1 µl SacI, 1 µl EcoRI, and 6 µl H2O. Digested DNA samples were electrophoresed overnight at 40 mA in a 0.85% SeaKem LE agarose gel (FMC, Rockland, Maine). The gel was blotted onto Millipore Immobilon-Ny+ (Bedford, MA) membrane overnight in 20X SSC (NaCl 175.2 g/l, Na citrate 88 g/l). The probe DNA was cut with BamHI/SacI (NEB, Beverly, MA) from pDAB1551 plasmid, which released a 7356 bp fragment containing the open

25 reading frame of the rebuilt *tcdA* gene. This 7356 bp fragment was labeled with P32 using a Stratagene Prime-it

RmT dCTP-Labeling Reactions kit (La Jolla, CA) and used for Southern hybridization. Hybridization was conducted in hybridization buffer (10% polyethylene glycol, 7% SDS [Sodium dodecyl sulfate], 0.6X SSC, 10 mM NaPO₄, 5 mM 5 EDTA, 10 µg/ml denatured salmon sperm) at 60 °C overnight. After hybridization, the membrane was washed with 10X SSC plus 0.1% SDS at 60 °C for 30 min and exposed to X ray film (Hyperfilm® MP, Amersham Life Sciences, Piscataway, NJ) for 1-2 days.

10

Results summarized indicate that a pattern of 8 hybridizing bands (the size of the expected fragment and larger) cosegregated with protein expression in 50% of all progeny assayed. These results are characteristic of 15 a complex insertion at a single site. All seedlings containing the insert also expressed toxin protein.

Example 6
Transformation Of Rice With a Vector Carrying Plasmid
20 pDAB1553 Encoding *Photorhabdus* Toxins

A. Plasmid pDAB1553

Plasmid pDAB1553 containing *tcdA* driven by the maize ubiquitin1 promoter and *hpt* (hygromycin 25 phosphotransferase providing resistance to the antibiotic hygromycin) under the control of 35T (a modified 35S promoter), was used for transformation.

Preparation of rice transformation vectors was 30 accomplished in two steps. First, a modified plant-optimized *tcdA* coding region was ligated into a rice plant expression cassette plasmid. In this step, the coding region was placed under the transcriptional control of a promoter functional in plant cells. RNA 35 transcription termination and polyadenylation were mediated by a downstream copy of the terminator region from the *Agrobacterium* nopaline synthase gene. One

plasmid designed to function in this role is plasmid pDAB1538 (described in the section on maize transformation vectors). In the second step, the complete gene comprised of the promoter, coding region, 5 and terminator region was ligated to a rice plant transformation vector that contained a plant expressible selectable marker gene which allowed the selection of transformed rice plant cells amongst a background of nontransformed cells. An example of such a vector is 10 pDAB354-Not1.

It is a feature of pDAB354-Not1 that the hygromycin phosphotransferase protein, which has as its substrate hygromycin B and related compounds, is produced in plant cells through transcription of its coding region mediated 15 by the Cauliflower Mosaic Virus 35S promoter and that termination of transcription plus polyadenylation are mediated by the nopaline synthase terminator region. It is further a feature of pDAB354-Not1 that any DNA fragment containing flanking NotI sites can be cloned 20 into the unique NotI site of pDAB354-Not1, thus physically linking the introduced DNA fragment to the aforementioned selectable marker gene.

To prepare a plant-expressible gene to produce the non-targeted TcdA protein in rice plant cells, DNA of a 25 plasmid (pAOH_4-OPTI) containing the plant-optimized *tcdA* coding region, (SEQ ID No:3) was cleaved with restriction enzymes *Nco*I and *Sac*I, and the large 7550 bp fragment was ligated to similarly-cut DNA of plasmid pDAB1538 to produce plasmid pDAB1551. DNA of pDAB1551 was then 30 digested with NotI, and the large 9933 bp fragment was ligated to NotI digested DNA of pDAB354-Not1 to produce plasmid pDAB1553.

It is a feature of plasmid pDAB1553 that the ubi1 and 35S promoters are encoded on the same DNA strand.

35 B. Production of Rice transgenics

For initiation of embryogenic callus, mature seeds of a *Japonica* cultivar, Taipei 309 were dehusked and surface-sterilized in 70% ethanol for 2-5 min. followed by a 30-45 min soak in 50% commercial bleach (2.6% sodium hypochlorite) with a few drops of 'Liquinox' soap. The seeds were then rinsed 3 times in sterile distilled water and placed on filter paper before transferring to 'callus induction' medium (i.e., NB). The NB medium consisted of N6 macro elements (Chu, 1978, The N6 medium and its application to anther culture of cereal crops. Proc. Symp. Plant Tissue Culture, Peking Press, p43-56), B5 micro elements and vitamins (Gamborg et al., 1968, Nutrient requirements of suspension cultures of soybean root cells. Exp. Cell Res. 50: 151-158), 300 mg/L casein hydrolysate, 500 mg/L L-proline, 500 mg/L L-glutamine, 30 g/L sucrose, 2 mg/L 2,4-dichloro-phenoxyacetic acid (2,4-D), and 2.5 g/L gelrite (Schweizerhall, NJ) with the pH adjusted to 5.8. The mature seed cultured on 'induction' media were incubated in the dark at 28°C. After 3 weeks of culture, the emerging primary callus induced from the scutellar region of mature embryo was transferred to fresh NB medium for further maintenance.

About 140 µg of plasmid pDAB1553 DNA was precipitated onto 60 mg of 1.0 micron (Bio-Rad) gold particles as described herein.

For helium blasting, actively growing embryogenic callus cultures, 2-4 mm in size, were subjected to a high osmoticum treatment. This treatment included placing of callus on NB medium with 0.2 M mannitol and 0.2 M sorbitol (Vain et al., 1993, Osmoticum treatment enhances particle bombardment-mediated transient and stable transformation of maize. Plant Cell Rep. 12: 84-88) for 4 h before helium blasting. Following osmoticum treatment, callus cultures were transferred to 'blasting' medium (NB+2% agar) and covered with a stainless steel screen (230 micron). The callus cultures were blasted at

2,000 psi helium pressures twice per target. After blasting, callus was transferred back to the media with high osmoticum overnight before placing on selection medium, which consisted NB medium with 30 mg/L hygromycin. After 2 weeks, the cultures were transferred to fresh selection medium with a higher concentration of selection agent, i.e., NB+50mg/L hygromycin (Li et al., 1993, An improved rice transformation system using the biolistic method. Plant Cell Rep. 12: 250-255).

10 Compact, white-yellow, embryogenic callus cultures, recovered on NB+50 mg/L hygromycin, were regenerated by transferring to 'pre-regeneration' (PR) medium + 50 mg/L hygromycin. The PR medium consisted of NB medium with 2 mg/L benzyl aminopurine (BAP), 1 mg/L naphthalene acetic acid (NAA), and 5 mg/L abscisic acid (ABA). After 2 weeks of culture in the dark, they were transferred to 'regeneration' (RN) medium. The composition of RN medium is NB medium with 3 mg/L BAP, and 0.5 mg/L NAA. The cultures on RN medium were incubated for 2 weeks at 20 28° C under high fluorescent light (325-ft-candles). The plantlets with 2 cm shoot were transferred to 1/2 MS medium (Murashige and Skoog, 1962, A revised medium for rapid growth and bioassays with tobacco tissue cultures. Physiol. Plant.15:473-497) with 1/2 B5 vitamins, 10 g/L sucrose, 0.05 mg/L NAA, 50 mg/L hygromycin and 2.5 g/L gelrite adjusted to pH 5.8 in magenta boxes. When plantlets were established with well-developed root systems, they were transferred to soil (1 metromix: 1 top soil) and raised in the greenhouse (29/24°C day/night cycle, 50-60% humidity, 12 h photoperiod) until maturity.

EXAMPLE 7

Chacterization Of Transgenic Rice Plants Expressing
35 Photorhabdus Toxin That Confer Insect Control.

A. Insect bioassays

Insect bioassays were performed using leaf discs and shown to be highly effective in controlling Southern corn rootworm. *Diabrotica undecimpunctata howardi* eggs are obtained from French Ag Research and hatched in petri dishes held at 28.5°C and 40% RH. The aerial parts are sampled from the transgenic plants and placed, singly into inverted petri dishes (100x15mm) containing 15ml of 1.6% aqueous agar in the bottom to provide humidity and filter paper in the top to absorb condensation. These preparations are infested with five neonate larvae per dish and held at 28.5°C and 40% RH for 3 days. Mortality and larval weights are recorded. Weight data were transformed using a logarithmic function to correct a correlation between the magnitude of the mean and variance.

Table 11

Treatment	Average Survivor Weight in mg ¹ (Duncan's Grouping)	Presence TcdA greenhouse-grown plants (number of +/number of plants tested)
GUS Control	0.390 A	-
1553-33	0.170 BCD	++
1553-44	0.167 BCD	+++
1553-62	0.125 CD	+++
1553-41	0.100 D	+++

Note: Means followed by the same letter are not significantly different based on Duncan's multiple range test (alpha=0.05).

Insect groups weighing less than 0.1 mg were set to 0.03 mg instead of zero to conduct a more conservative analysis. Weight data were transformed (Log10) for analyses. A single replicate was used on each of three test dates. Plants were sampled from magenta boxes. The results demonstrate that in leaf disc bioassays, several rice events derived by transformation with *tcdA* gene were demonstrated to statistically have a functional affect on corn rootworm neonate.

Claims

1. An isolated nucleic acid of SEQ ID NO: 3 or SEQ ID NO:4.
2. A transgenic monocot cell having a genome comprising SEQ ID NO:3 or SEQ ID NO:4.
3. A transgenic dicot cell having a genome comprising SEQ ID NO:3 or SEQ ID NO:4.
4. A transgenic plant with a genome comprising a nucleic acid of SEQ ID NO: 3 or SEQ ID NO:4 that imparts insect resistance.
5. A transgenic plant of claim 4 wherein the plant is rice.
6. A transgenic plant of claim 4 wherein the plant is maize.
- 15 7. A transgenic plant of claim 4 wherein the plant is tobacco.

SEQUENCE LISTING

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<120> Transgenic Plants Expressing *Photorhabdus* Toxin

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<150> US 60/148,356
 <151> 1999-08-11

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<170> PatentIn Ver. 2.0

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<221> CDS

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cgc cag caa gta tct gag cac ctc tcc tgg tcc gaa aca cac gac tta 144
 Arg Gln Gln Val Ser Glu His Leu Ser Trp Ser Glu Thr His Asp Leu
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tat cat gat gca caa cag gca caa aag gat aat cgc ctg tat gaa gcg 192
 Tyr His Asp Ala Gln Gln Ala Gln Lys Asp Asn Arg Leu Tyr Glu Ala
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cgt att ctc aaa cgc gcc aat ccc caa tta caa aat gcg gtg cat ctt 240
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gcc att ctc gct ccc aat gct gaa ctg ata ggc tat aac aat caa ttt 288
 Ala Ile Leu Ala Pro Asn Ala Glu Leu Ile Gly Tyr Asn Asn Gln Phe
 85 90 95

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 Ser Gly Arg Ala Ser Gln Tyr Val Ala Pro Gly Thr Val Ser Ser Met

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tta cac gca agt gac tcc gtt tat tat ctg gat acc cgc cgc cca gat Leu His Ala Ser Asp Ser Val Tyr Tyr Leu Asp Thr Arg Arg Pro Asp 130	135	140	432
ctc aaa tca atg gcg ctc agt cag caa aat atg gat ata gaa tta tcc Leu Lys Ser Met Ala Leu Ser Gln Gln Asn Met Asp Ile Glu Leu Ser 145	150	155	480
aca ctc tct ttg tcc aat gag ctg tta ttg gaa agc att aaa act gaa Thr Leu Ser Leu Ser Asn Glu Leu Leu Leu Glu Ser Ile Lys Thr Glu 165	170	175	528
tct aaa ctg gaa aac tat act aaa gtg atg gaa atg ctc tcc act ttc Ser Lys Leu Glu Asn Tyr Thr Lys Val Met Glu Met Leu Ser Thr Phe 180	185	190	576
cgt cct tcc ggc gca acg cct tat cat gat gct tat gaa aat gtg cgt Arg Pro Ser Gly Ala Thr Pro Tyr His Asp Ala Tyr Glu Asn Val Arg 195	200	205	624
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Tyr	Arg	Leu	Asp	Tyr	Lys	Phe	Lys	Asn	Phe	Tyr	Asn	Ala	Ser	Tyr	Leu	
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								360						365		
tcc	atc	aag	tta	aat	gat	aaa	aga	gaa	ctt	gtt	cga	act	gaa	ggc	gct	1152
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Pro	Gln	Val	Asn	Ile	Glu	Tyr	Ser	Ala	Asn	Ile	Thr	Leu	Asn	Thr	Ala	
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gat	atc	agt	caa	cct	ttt	gaa	att	ggc	ctg	aca	cga	gta	ctt	cct	tcc	1248
Asp	Ile	Ser	Gln	Pro	Phe	Glu	Ile	Gly	Leu	Thr	Arg	Val	Leu	Pro	Ser	
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Gly	Ser	Trp	Ala	Tyr	Ala	Ala	Lys	Phe	Thr	Val	Glu	Glu	Tyr	Asn		
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caa	tac	tct	ttt	ctg	cta	aaa	ctt	aac	aag	gct	att	cgt	cta	tca	cgt	1344
Gln	Tyr	Ser	Phe	Leu	Leu	Lys	Leu	Asn	Lys	Ala	Ile	Arg	Leu	Ser	Arg	
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Ile	Leu	Cys	Asn	Ala	Pro	Ile	Ser	Gln	Arg	Ser	Tyr	Asp	Asn	Gln	Pro	
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Phe	Ser	Thr	Gly	Asp	Glu	Glu	Ile	Asp	Leu	Asn	Ser	Gly	Ser	Thr	Gly	
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gat	tgg	cga	aaa	acc	ata	ctt	aag	cgt	gca	ttt	aat	att	gat	gat	gtc	1680
Asp	Trp	Arg	Lys	Thr	Ile	Leu	Lys	Arg	Ala	Phe	Asn	Ile	Asp	Asp	Val	
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Pro Tyr Lys Ser Thr Ile Arg Pro Val Ile Tyr Lys Ser Arg Leu Tyr			
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 1380 1385 1390

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 Val Asn Pro Asn Asn Ser Ser Asn Lys Leu Met Phe Tyr Pro Val Tyr
 1395 1400 1405

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 Gly Gly Lys Glu Gln Thr Phe Thr Ala Asp Lys Asp Val Ser Ile Gln
 1505 1510 1515 1520

cca tca cct agc ttt gat gaa atg aat tat caa ttt aat gcc ctt gaa 4608
 Pro Ser Pro Ser Phe Asp Glu Met Asn Tyr Gln Phe Asn Ala Leu Glu
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ata gac ggt tct ggt ctg aat ttt att aac aac tca gcc agt att gat 4656
 Ile Asp Gly Ser Gly Leu Asn Phe Ile Asn Asn Ser Ala Ser Ile Asp
 1540 1545 1550

gtt act ttt acc gca ttt gcg gag gat ggc cgc aaa ctg ggt tat gaa 4704
 Val Thr Phe Thr Ala Phe Ala Glu Asp Gly Arg Lys Leu Gly Tyr Glu
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agt ttc agt att cct gtt acc ctc aag gta agt acc gat aat gcc ctg 4752
 Ser Phe Ser Ile Pro Val Thr Leu Lys Val Ser Thr Asp Asn Ala Leu
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acc ctg cac cat aat gaa aat ggt gcg caa tat atg caa tgg caa tcc 4800
 Thr Leu His His Asn Glu Asn Gly Ala Gln Tyr Met Gln Trp Gln Ser
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tat cgt acc cgc ctg aat act cta ttt gcc cgc cag tgg gtt gca cgc 4848
 Tyr,Arg Thr Arg Leu Asn Thr Leu Phe Ala Arg Gln Leu Val Ala Arg
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gcc acc acc gga atc gat aca att ctg agt atg gaa act cag aat att 4896
 Ala Thr Thr Gly Ile Asp Thr Ile Leu Ser Met Glu Thr Gln Asn Ile
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 Gln Glu Pro Gln Leu Gly Lys Gly Phe Tyr Ala Thr Phe Val Ile Pro
 1635 1640 1645

ccc tat aac cta tca act cat ggt gat gaa cgt tgg ttt aag ctt tat 4992
 Pro Tyr Asn Leu Ser Thr His Gly Asp Glu Arg Trp Phe Lys Leu Tyr
 1650 1655 1660

atc aaa cat gtt gtt gat aat aat tca cat att atc tat tca ggc cag 5040
 Ile Lys His Val Val Asp Asn Asn Ser His Ile Ile Tyr Ser Gly Gln
 1665 1670 1675 1680

cta aca gat aca aat ata aac atc aca tta ttt att cct ctt gat gat 5088
 Leu Thr Asp Thr Asn Ile Asn Ile Thr Leu Phe Ile Pro Leu Asp Asp
 1685 1690 1695

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 Val Pro Leu Asn Gln Asp Tyr His Ala Lys Val Tyr Met Thr Phe Lys
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 Lys Ser Pro Ser Asp Gly Thr Trp Trp Gly Pro His Phe Val Arg Asp
 1715 1720 1725

gat aaa gga ata gta aca ata aac cct aaa tcc att ttg acc cat ttt 5232
 Asp Lys Gly Ile Val Thr Ile Asn Pro Lys Ser Ile Leu Thr His Phe
 1730 1735 1740

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 Glu Ser Val Asn Val Leu Asn Asn Ile Ser Ser Glu Pro Met Asp Phe
 1745 1750 1755 1760

agc ggc gct aac agc ctc tat ttc tgg gaa ctg ttc tac tat acc ccg 5328
 Ser Gly Ala Asn Ser Leu Tyr Phe Trp Glu Leu Phe Tyr Tyr Thr Pro
 1765 1770 1775

atg ctg gtt gct caa cgt ttg ctg cat gaa cag aac ttc gat gaa gcc 5376
 Met Leu Val Ala Gln Arg Leu Leu His Glu Gln Asn Phe Asp Glu Ala
 1780 1785 1790

aac cgt tgg ctg aaa tat gtc tgg agt cca tcc ggt tat att gtc cac 5424
 Asn Arg Trp Leu Lys Tyr Val Trp Ser Pro Ser Gly Tyr Ile Val His
 1795 1800 1805

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Gly Gln Ile Gln Asn Tyr Gln Trp Asn Val Arg Pro Leu Leu Glu Asp
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 Thr Ser Trp Asn Ser Asp Pro Leu Asp Ser Val Asp Pro Asp Ala Val
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gca cag cac gat cca atg cac tac aaa gtt tca act ttt atg cgt acc 5568
 Ala Gln His Asp Pro Met His Tyr Lys Val Ser Thr Phe Met Arg Thr
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 Leu Asp Leu Leu Ile Ala Arg Gly Asp His Ala Tyr Arg Gln Leu Glu
 1860 1865 1870

cga gat aca ctc aac gaa gcg aag atg tgg tat atg caa gcg ctg cat 5664
 Arg Asp Thr Leu Asn Glu Ala Lys Met Trp Tyr Met Gln Ala Leu His
 1875 1880 1885

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 Leu Leu Gly Asp Lys Pro Tyr Leu Pro Leu Ser Thr Thr Trp Ser Asp
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 Pro Arg Leu Asp Arg Ala Ala Asp Ile Thr Thr Gln Asn Ala His Asp
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 Ser Ala Ile Val Ala Leu Arg Gln Asn Ile Pro Thr Pro Ala Pro Leu
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tca ttg cgc agc gct aat acc ctg act gat ctc ttc ctg ccg caa atc 5856
 Ser Leu Arg Ser Ala Asn Thr Leu Thr Asp Leu Phe Leu Pro Gln Ile
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 Asn Glu Val Met Met Asn Tyr Trp Gln Thr Leu Ala Gln Arg Val Tyr
 1955 1960 1965

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 Asn Leu Arg His Asn Leu Ser Ile Asp Gly Gln Pro Leu Tyr Leu Pro
 1970 1975 1980

atc tat gcc aca ccg gcc gat ccg aaa gcg tta ctc agc gcc gcc gtt 6000
 Ile Tyr Ala Thr Pro Ala Asp Pro Lys Ala Leu Leu Ser Ala Ala Val
 1985 1990 1995 2000

gcc act tct caa ggt gga ggc aag cta ccg gaa tca ttt atg tcc ctg 6048
 Ala Thr Ser Gln Gly Gly Lys Leu Pro Glu Ser Phe Met Ser Leu
 2005 2010 2015

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 Trp Arg Phe Pro His Met Leu Glu Asn Ala Arg Gly Met Val Ser Gln
 2020 2025 2030

ctc acc cag ttc ggc tcc acg tta caa aat att atc gaa cgt cag gac 6144
 Leu Thr Gln Phe Gly Ser Thr Leu Gln Asn Ile Ile Glu Arg Gln Asp
 2035 2040 2045

gcg gaa gcg ctc aat gcg tta tta caa aat cag gcc ggc gag ctg ata 6192
 Ala Glu Ala Leu Asn Ala Leu Leu Gln Asn Gln Ala Ala Glu Leu Ile

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ttg act aac ctg agc att cag gac aaa acc att gaa gaa ttg gat gcc Leu Thr Asn Leu Ser Ile Gln Asp Lys Thr Ile Glu Glu Leu Asp Ala	2065	2070	6240
2075			2080
gag aaa acg gtg ttg gaa aaa tcc aaa gcg gga gca caa tcg cgc ttt Glu Lys Thr Val Leu Glu Lys Ser Lys Ala Gly Ala Gln Ser Arg Phe	2085	2090	6288
2095			
gat agc tac ggc aaa ctg tac gat gag aat atc aac gcc ggt gaa aac Asp Ser Tyr Gly Lys Leu Tyr Asp Glu Asn Ile Asn Ala Gly Glu Asn	2100	2105	6336
2110			
caa gcc atg acg cta cga gcg tcc gcc ggg ctt acc acg gca gtt Gln Ala Met Thr Leu Arg Ala Ser Ala Ala Gly Leu Thr Thr Ala Val	2115	2120	6384
2125			
cag gca tcc cgt ctg gcc ggt gcg gcg gct gat ctg gtg cct aac atc Gln Ala Ser Arg Leu Ala Gly Ala Ala Asp Leu Val Pro Asn Ile	2130	2135	6432
2140			
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2155			2160
aca ggt tat gtg atg gaa ttc tcc gcg aat gtt atg aac acc gaa gcg Thr Gly Tyr Val Met Glu Phe Ser Ala Asn Val Met Asn Thr Glu Ala	2165	2170	6528
2175			
gat aaa att agc caa tct gaa acc tac cgt cgt cgc cgt cag gag tgg Asp Lys Ile Ser Gln Ser Glu Thr Tyr Arg Arg Arg Arg Gln Glu Trp	2180	2185	6576
2190			
gag atc cag cgg aat aat gcc gaa gcg gaa ttg aag caa atc gat gct Glu Ile Gln Arg Asn Asn Ala Glu Ala Glu Leu Lys Gln Ile Asp Ala	2195	2200	6624
2205			
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2220			
acc agt ctg aaa acc caa caa gaa cag acc caa tct caa ttg gcc ttc Thr Ser Leu Lys Thr Gln Gln Glu Gln Thr Gln Ser Gln Leu Ala Phe	2225	2230	6720
2235			2240
ctg caa cgt aag ttc agc aat cag gcg tta tac aac tgg ctg cgt ggt Leu Gln Arg Lys Phe Ser Asn Gln Ala Leu Tyr Asn Trp Leu Arg Gly	2245	2250	6768
2255			
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2270			
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2285			
gcc cgc ttc att aaa ccg ggc gcc tgg cag gga acc tat gcc ggt ctg Ala Arg Phe Ile Lys Pro Gly Ala Trp Gln Gly Thr Tyr Ala Gly Leu	2290	2295	6912
2300			

ctt gca ggt gaa acc ttg atg ctg agt ctg gca caa atg gaa gac gct Leu Ala Gly Glu Thr Leu Met Leu Ser Leu Ala Gln Met Glu Asp Ala 2305 2310 2315 2320	6960
cat ctg aaa cgc gat aaa cgc gca tta gag gtt gaa cgc aca gta tcg His Leu Lys Arg Asp Lys Arg Ala Leu Glu Val Glu Arg Thr Val Ser 2325 2330 2335	7008
ctg gcc gaa gtt tat gca gga tta cca aaa gat aac ggt cca ttt tcc Leu Ala Glu Val Tyr Ala Gly Leu Pro Lys Asp Asn Gly Pro Phe Ser 2340 2345 2350	7056
ctg gct cag gaa att gac aag ctg gtg agt caa ggt tca ggc agt gcc Leu Ala Gln Glu Ile Asp Lys Leu Val Ser Gln Gly Ser Gly Ser Ala 2355 2360 2365	7104
ggc agt ggt aat aat att ttg gcg ttc ggc gcc ggc acg gac act aaa Gly Ser Gly Asn Asn Asn Leu Ala Phe Gly Ala Gly Thr Asp Thr Lys 2370 2375 2380	7152
acc tct ttg cag gca tca gtt tca ttc gct gat ttg aaa att cgt gaa Thr Ser Leu Gln Ala Ser Val Phe Ala Asp Leu Lys Ile Arg Glu 2385 2390 2395 2400	7200
gat tac ccg gca tcg ctt ggc aaa att cga cgt atc aaa cag atc agc Asp Tyr Pro Ala Ser Leu Gly Lys Ile Arg Arg Ile Lys Gln Ile Ser 2405 2410 2415	7248
gtc act ttg ccc gcg cta ctg gga ccg tat cag gat gta cag gca ata Val Thr Leu Pro Ala Leu Leu Gly Pro Tyr Gln Asp Val Gln Ala Ile 2420 2425 2430	7296
ttg tct tac ggc gat aaa gcc gga tta gct aac ggc tgt gaa ggc ctg Leu Ser Tyr Gly Asp Lys Ala Gly Leu Ala Asn Gly Cys Glu Ala Leu 2435 2440 2445	7344
gca gtt tct cac ggt atg aat gac agc ggc caa ttc cag ctc gat ttc Ala Val Ser His Gly Met Asn Asp Ser Gly Gln Phe Gln Leu Asp Phe 2450 2455 2460	7392
aac gat ggc aaa ttc ctg cca ttc gaa ggc atc gcc att gat caa ggc Asn Asp Gly Lys Phe Leu Pro Phe Glu Gly Ile Ala Ile Asp Gln Gly 2465 2470 2475 2480	7440
acg ctg aca ctg agc ttc cca aat gca tct atg ccg gag aaa ggt aaa Thr Leu Thr Leu Ser Phe Pro Asn Ala Ser Met Pro Glu Lys Gly Lys 2485 2490 2495	7488
caa gcc act atg tta aaa acc ctg aac gat atc att ttg cat att cgc Gln Ala Thr Met Leu Lys Thr Leu Asn Asp Ile Ile Leu His Ile Arg 2500 2505 2510	7536
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caa tta act tgt ccg gcg gaa att gct ttg tat ccc ttt gat act ttc	96
Gln Leu Thr Cys Pro Ala Glu Ile Ala Leu Tyr Pro Phe Asp Thr Phe	
20 25 30	

cgg gaa aaa act cgg gga atg gtt aat tgg ggg gaa gca aaa cgg att	144
Arg Glu Lys Thr Arg Gly Met Val Asn Trp Gly Glu Ala Lys Arg Ile	
35 40 45	

tat gaa att gca caa gcg gaa cag gat aga aac cta ctt cat gaa aaa	192
Tyr Glu Ile Ala Gln Ala Glu Gln Asp Arg Asn Leu Leu His Glu Lys	
50 55 60	

cgt att ttt gcc tat gct aat ccg ctg ctg aaa aac gct gtt cgg ttg	240
Arg Ile Phe Ala Tyr Ala Asn Pro Leu Leu Lys Asn Ala Val Arg Leu	
65 70 75 80	

ggt acc cgg caa atg ttg ggt ttt ata caa ggt tat agt gat ctg ttt	288
Gly Thr Arg Gln Met Leu Gly Phe Ile Gln Gly Tyr Ser Asp Leu Phe	
85 90 95	

ggt aat cgt gct gat aac tat gcc gcg ccg ggc tcg gtt gca tcg atg	336
Gly Asn Arg Ala Asp Asn Tyr Ala Ala Pro Gly Ser Val Ala Ser Met	
100 105 110	

ttc tca ccg gcg gct tat ttg acg gaa ttg tac cgt gaa gcc aaa aac	384
Phe Ser Pro Ala Ala Tyr Leu Thr Glu Leu Tyr Arg Glu Ala Lys Asn	
115 120 125	

ttg cat gac agc agc tca att tat tac cta gat aaa cgt cgc ccg gat	432
Leu His Asp Ser Ser Ser Ile Tyr Tyr Leu Asp Lys Arg Arg Pro Asp	
130 135 140	

tta gca agc tta atg ctc agc cag aaa aat atg gat gag gaa att tca	480
Leu Ala Ser Leu Met Leu Ser Gln Lys Asn Met Asp Glu Glu Ile Ser	
145 150 155 160	

acg ctg gct ctc tct aat gaa ttg tgc ctt gcc ggg atc gaa aca aaa	528
Thr Leu Ala Leu Ser Asn Glu Leu Cys Leu Ala Gly Ile Glu Thr Lys	
165 170 175	

aca gga aaa tca caa gat gaa gtg atg gat atg ttg tca act tat cgt	576
Thr Gly Lys Ser Gln Asp Glu Val Met Asp Met Leu Ser Thr Tyr Arg	
180 185 190	

tta agt gga gag aca cct tat cat cac gct tat gaa act gtt cgt gaa	624
Leu Ser Gly Glu Thr Pro Tyr His His Ala Tyr Glu Thr Val Arg Glu	
195 200 205	

atc gtt cat gaa cgt gat cca gga ttt cgt cat ttg tca cag gca ccc Ile Val His Glu Arg Asp Pro Gly Phe Arg His Leu Ser Gln Ala Pro 210	215	220	672	
att gtt gct gct aag ctc gat cct gtg act ttg ttg ggt att agc tcc Ile Val Ala Ala Lys Leu Asp Pro Val Thr Leu Leu Gly Ile Ser Ser 225	230	235	240	720
cat att tcg cca gaa ctg tat aac ttg ctg att gag gag atc ccg gaa His Ile Ser Pro Glu Leu Tyr Asn Leu Leu Ile Glu Glu Ile Pro Glu 245	250	255	255	768
aaa gat gaa gcc gcg ctt gat acg ctt tat aaa aca aac ttt ggc gat Lys Asp Glu Ala Ala Leu Asp Thr Leu Tyr Lys Thr Asn Phe Gly Asp 260	265	270	270	816
att act act gct tta atg tcc cca agt tat ctg gcc ccg tat tat Ile Thr Thr Ala Gln Leu Met Ser Pro Ser Tyr Leu Ala Arg Tyr Tyr 275	280	285	285	864
ggc gtc tca ccg gaa gat att gcc tac gtg acg act tca tta tca cat Gly Val Ser Pro Glu Asp Ile Ala Tyr Val Thr Thr Ser Leu Ser His 290	295	300	300	912
gtt gga tat agc agt gat att ctg gtt att ccg ttg gtc gat ggt gtg Val Gly Tyr Ser Ser Asp Ile Leu Val Ile Pro Leu Val Asp Gly Val 305	310	315	320	960
ggt aag atg gaa gta gtt cgt gtt acc cga aca cca tcg gat aat tat Gly Lys Met Glu Val Val Arg Val Thr Arg Thr Pro Ser Asp Asn Tyr 325	330	335	335	1008
acc agt cag acg aat tat att gag ctg tat cca cag ggt ggc gac aat Thr Ser Gln Thr Asn Tyr Ile Glu Leu Tyr Pro Gln Gly Asp Asn 340	345	350	350	1056
tat ttg atc aaa tac aat cta agc aat agt ttt ggt ttg gat gat tat Tyr Leu Ile Lys Tyr Asn Leu Ser Asn Ser Phe Gly Leu Asp Asp Phe 355	360	365	365	1104
tat ctg caa tat aaa gat ggt tcc gct gat tgg act gag att gcc cat Tyr Leu Gln Tyr Lys Asp Gly Ser Ala Asp Trp Thr Glu Ile Ala His 370	375	380	380	1152
aat ccc tat cct gat atg gtc ata aat caa aag tat gaa tca cag gcg Asn Pro Tyr Pro Asp Met Val Ile Asn Gln Lys Tyr Glu Ser Gln Ala 385	390	395	400	1200
aca atc aaa cgt agt gac tct gac aat ata ctc agt ata ggg tta caa Thr Ile Lys Arg Ser Asp Ser Asp Asn Ile Leu Ser Ile Gly Leu Gln 405	410	415	415	1248
aga tgg cat agc ggt agt tat aat ttt gcc gcc gcc aat ttt aaa att Arg Trp His Ser Gly Ser Tyr Asn Phe Ala Ala Asn Phe Lys Ile 420	425	430	430	1296
gac caa tac tcc ccg aaa gct ttc ctg ctt aaa atg aat aag gct att Asp Gln Tyr Ser Pro Lys Ala Phe Leu Leu Lys Met Asn Lys Ala Ile 435	440	445	445	1344
cgg ttg ctc aaa gct acc ggc ctc tct ttt gct acg ttg gag cgt att				1392

Arg Leu Leu Lys Ala Thr Gly Leu Ser Phe Ala Thr Leu Glu Arg Ile		
450	455	460
gtt gat agt gtt aat agc acc aaa tcc atc acg gtt gag gta tta aac		1440
Val Asp Ser Val Asn Ser Thr Lys Ser Ile Thr Val Glu Val Leu Asn		
465	470	475
480		
aag gtt tat cgg gta aaa ttc tat att gat cgt tat ggc atc agt gaa		1488
Lys Val Tyr Arg Val Lys Phe Tyr Ile Asp Arg Tyr Gly Ile Ser Glu		
485	490	495
gag aca gcc gct att ttg gct aat att aat atc tct cag caa gct gtt		1536
Glu Thr Ala Ala Ile Leu Ala Asn Ile Asn Ile Ser Gln Gln Ala Val		
500	505	510
ggc aat cag ctt agc cag ttt gag caa cta ttt aat cac ccc ccc ctc		1584
Gly Asn Gln Leu Ser Gln Phe Glu Gln Leu Phe Asn His Pro Pro Leu		
515	520	525
aat ggt att cgc tat gaa atc agt gag gac aac tcc aaa cat ctt cct		1632
Asn Gly Ile Arg Tyr Glu Ile Ser Glu Asp Asn Ser Lys His Leu Pro		
530	535	540
aat cct gat ctg aac ctt aaa cca gac agt acc ggt gat gat caa cgc		1680
Asn Pro Asp Leu Asn Leu Lys Pro Asp Ser Thr Gly Asp Asp Gln Arg		
545	550	555
560		
aag gcg gtt tta aaa cgc gcg ttt cag gtt aac gcc agt gag ttg tat		1728
Lys Ala Val Leu Lys Arg Ala Phe Gln Val Asn Ala Ser Glu Leu Tyr		
565	570	575
cag atg tta ttg atc act gat cgt aaa gaa gac ggt gtt atc aaa aat		1776
Gln Met Leu Leu Ile Thr Asp Arg Lys Glu Asp Gly Val Ile Lys Asn		
580	585	590
aac tta gag aat ttg tct gat ctg tat ttg gtt agt ttg ctg gcc cag		1824
Asn Leu Glu Asn Leu Ser Asp Leu Tyr Leu Val Ser Leu Leu Ala Gln		
595	600	605
att cat aac ctg act att gct gaa ttg aac att ttg ttg gtg att tgt		1872
Ile His Asn Leu Thr Ile Ala Glu Leu Asn Ile Leu Leu Val Ile Cys		
610	615	620
ggc tat ggc gac acc aac att tat cag att acc gac gat aat tta gcc		1920
Gly Tyr Gly Asp Thr Asn Ile Tyr Gln Ile Thr Asp Asp Asn Leu Ala		
625	630	635
640		
aaa ata gtg gaa aca ttg ttg tgg atc act caa tgg ttg aag acc caa		1968
Lys Ile Val Glu Thr Leu Leu Trp Ile Thr Gln Trp Leu Lys Thr Gln		
645	650	655
aaa tgg aca gtt acc gac ctg ttt ctg atg acc acg gcc act tac agc		2016
Lys Trp Thr Val Thr Asp Leu Phe Leu Met Thr Thr Ala Thr Tyr Ser		
660	665	670
acc act tta acg cca gaa att agc aat ctg acg gct acg ttg tct tca		2064
Thr Thr Leu Thr Pro Glu Ile Ser Asn Leu Thr Ala Thr Leu Ser Ser		
675	680	685
act ttg cat ggc aaa gag agt ctg att ggg gaa gat ctg aaa aga gca		2112
Thr Leu His Gly Lys Glu Ser Leu Ile Gly Glu Asp Leu Lys Arg Ala		

690	695	700														
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Met	Ala	Pro	Cys	Phe	Thr	Ser	Ala	Leu	His	Leu	Thr	Ser	Gln	Glu	Val	
705							710			715				720		
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Ala	Tyr	Asp	Leu	Leu	Leu	Trp	Ile	Asp	Gln	Ile	Gln	Pro	Ala	Gln	Ile	
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Thr	Val	Asp	Gly	Phe	Trp	Glu	Glu	Val	Gln	Thr	Thr	Pro	Thr	Ser	Leu	
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Lys	Val	Ile	Thr	Phe	Ala	Gln	Val	Leu	Ala	Gln	Leu	Ser	Leu	Ile	Tyr	
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cgt	cgt	att	ggg	tta	agt	gaa	acg	gaa	ctg	tca	ctg	atc	gtg	act	caa	2352
Arg	Arg	Ile	Gly	Leu	Ser	Glu	Thr	Glu	Leu	Ser	Leu	Ile	Val	Thr	Gln	
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tct	tct	ctg	cta	gtg	gca	ggc	aaa	agc	ata	ctg	gat	cac	ggt	ctg	tta	2400
Ser	Ser	Leu	Leu	Val	Ala	Gly	Lys	Ser	Ile	Leu	Asp	His	Gly	Leu	Leu	
							785			790				795		
acc	ctg	atg	gcc	ttg	gaa	ggt	ttt	cat	acc	tgg	gtt	aat	ggc	ttg	ggg	2448
Thr	Leu	Met	Ala	Leu	Glu	Gly	Phe	His	Thr	Trp	Val	Asn	Gly	Leu	Gly	
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caa	cat	gcc	tcc	ttg	ata	ttg	gct	gct	ttg	aaa	gac	gga	gcc	ttg	aca	2496
Gln	His	Ala	Ser	Leu	Ile	Leu	Ala	Ala	Leu	Lys	Asp	Gly	Ala	Leu	Thr	
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gtt	acc	gat	gta	gca	caa	gct	atg	aat	aag	gag	gaa	tct	ctc	cta	caa	2544
Val	Thr	Asp	Val	Ala	Gln	Ala	Met	Asn	Lys	Glu	Glu	Ser	Leu	Leu	Gln	
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Met	Ala	Ala	Asn	Gln	Val	Glu	Lys	Asp	Leu	Thr	Lys	Leu	Thr	Ser	Trp	
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aca	cag	att	gac	gct	att	ctg	caa	tgg	tta	cag	atg	tct	tcg	gcc	ttg	2640
Thr	Gln	Ile	Asp	Ala	Ile	Leu	Gln	Trp	Leu	Gln	Met	Ser	Ser	Ala	Leu	
							865			870				875		
gct	gtt	tct	cca	ctg	gat	ctg	gca	ggg	atg	atg	gct	gcc	ctg	aaa	tat	2688
Ala	Val	Ser	Pro	Leu	Asp	Leu	Ala	Gly	Met	Met	Ala	Leu	Lys	Tyr	Gly	
							885			890				895		
ata	gat	cat	aac	tat	gct	gcc	ttg	caa	gct	gct	gct	gct	ctg	atg	2736	
Ile	Asp	His	Asn	Tyr	Ala	Ala	Trp	Gln	Ala	Ala	Ala	Ala	Ala	Leu	Met	
							900			905				910		
gct	gat	cat	gct	aat	cag	gca	cag	aaa	aaa	ctg	gat	gag	acg	ttc	agt	2784
Ala	Asp	His	Ala	Asn	Gln	Ala	Gln	Lys	Lys	Leu	Asp	Glu	Thr	Phe	Ser	
							915			920				925		
aag	gca	tta	tgt	aac	tat	tat	att	aat	gct	gtt	gtc	gat	agt	gct	gct	2832
Lys	Ala	Leu	Cys	Asn	Tyr	Tyr	Ile	Asn	Ala	Val	Val	Asp	Ser	Ala	Ala	
							930			935				940		

gga gta cgt gat cgt aac ggt tta tat acc tat ttg ctg att gat aat	2880
Gly Val Arg Asp Arg Asn Gly Leu Tyr Thr Tyr Leu Leu Ile Asp Asn	
945 950 955 960	
cag gtt tct gcc gat gtg atc act tca cgt att gca gaa gct atc gcc	2928
Gln Val Ser Ala Asp Val Ile Thr Ser Arg Ile Ala Glu Ala Ile Ala	
965 970 975	
ggt att caa ctg tac gtt aac cgg gct tta aac cga gat gaa ggt cag	2976
Gly Ile Gln Leu Tyr Val Asn Arg Ala Leu Asn Arg Asp Glu Gly Gln	
980 985 990	
ctt gca tcg gac gtt agt acc cgt ca'g ttc ttc act gac tgg gaa cgt	3024
Leu Ala Ser Asp Val Ser Thr Arg Gln Phe Phe Thr Asp Trp Glu Arg	
995 1000 1005	
tac aat aaa cgt tac agt act tgg gct ggt gtc tct gaa ctg gtc tat	3072
Tyr Asn Lys Arg Tyr Ser Thr Trp Ala Gly Val Ser Glu Leu Val Tyr	
1010 1015 1020	
tat cca gaa aac tat gtt gat ccc act cag cgc att ggg caa acc aaa	3120
Tyr Pro Glu Asn Tyr Val Asp Pro Thr Gln Arg Ile Gly Gln Thr Lys	
1025 1030 1035 1040	
atg atg gat gcg ctg ttg caa tcc atc aac cag agc cag cta aat gcg	3168
Met Met Asp Ala Leu Leu Gln Ser Ile Asn Gln Ser Gln Leu Asn Ala	
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gat acg gtg gaa gat gct ttc aaa act tat ttg acc agc ttt gag cag	3216
Asp Thr Val Glu Asp Ala Phe Lys Thr Tyr Leu Thr Ser Phe Glu Gln	
1060 1065 1070	
gta gca aat ctg aaa gta att agt gct tac cac gat aat gtg aat gtg	3264
Val Ala Asn Leu Lys Val Ile Ser Ala Tyr His Asp Asn Val Asn Val	
1075 1080 1085	
gat caa gga tta act tat ttt atc ggt atc gac caa gca gct ccg ggt	3312
Asp Gln Gly Leu Thr Tyr Phe Ile Gly Ile Asp Gln Ala Ala Pro Gly	
1090 1095 1100	
acg tat tac tgg cgt agt gtt gat cac agc aaa tgt gaa aat ggc aag	3360
Thr Tyr Trp Arg Ser Val Asp His Ser Lys Cys Glu Asn Gly Lys	
1105 1110 1115 1120	
ttt gcc gct aat gct tgg ggt gag tgg aat aaa att acc tgt gct gtc	3408
Phe Ala Ala Asn Ala Trp Gly Glu Trp Asn Lys Ile Thr Cys Ala Val	
1125 1130 1135	
aat cct tgg aaa aat atc atc cgt ccg gtt tat atg tcc cgc tta	3456
Asn Pro Trp Lys Asn Ile Ile Arg Pro Val Val Tyr Met Ser Arg Leu	
1140 1145 1150	
tat ctg cta tgg ctg gag cag caa tca aag aaa agt gat gat ggt aaa	3504
Tyr Leu Leu Trp Leu Glu Gln Gln Ser Lys Lys Ser Asp Asp Gly Lys	
1155 1160 1165	
acc acg att tat caa tat aac tta aaa ctg gct cat att cgt tac gac	3552
Thr Thr Ile Tyr Gln Tyr Asn Leu Lys Leu Ala His Ile Arg Tyr Asp	
1170 1175 1180	

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 Gly Ser Trp Asn Thr Pro Phe Thr Phe Asp Val Thr Glu Lys Val Lys
 1185 1190 1195 1200

 aat tac acg tcg agt act gat gct gct gaa tct tta ggg ttg tat tgt 3648
 Asn Tyr Thr Ser Ser Thr Asp Ala Ala Glu Ser Leu Gly Leu Tyr Cys
 1205 1210 1215

 act ggt tat caa ggg gaa gac act cta tta gtt atg ttc tat tcg atg 3696
 Thr Gly Tyr Gln Gly Glu Asp Thr Leu Leu Val Met Phe Tyr Ser Met
 1220 1225 1230

 cag agt agt tat agc tcc tat acc gat aat aat gcg ccg gtc act ggg 3744
 Gln Ser Ser Tyr Ser Ser Tyr Asp Asn Asn Ala Pro Val Thr Gly
 1235 1240 1245

 cta tat att ttc gct gat atg tca tca gac aat atg acg aat gca caa 3792
 Leu Tyr Ile Phe Ala Asp Met Ser Ser Asp Asn Met Thr Asn Ala Gln
 1250 1255 1260

 gca act aac tat tgg aat aac agt tat ccg caa ttt gat act gtg atg 3840
 Ala Thr Asn Tyr Trp Asn Asn Ser Tyr Pro Gln Phe Asp Thr Val Met
 1265 1270 1275 1280

 gca gat ccg gat agc gac aat aaa aaa gtc ata acc aga aga gtt aat 3888
 Ala Asp Pro Asp Ser Asp Asn Lys Lys Val Ile Thr Arg Arg Val Asn
 1285 1290 1295

 aac cgt tat gcg gag gat tat gaa att cct tcc tct gtg aca agt aac 3936
 Asn Arg Tyr Ala Glu Asp Tyr Glu Ile Pro Ser Ser Val Thr Ser Asn
 1300 1305 1310

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 Ser Asn Tyr Ser Trp Gly Asp His Ser Leu Thr Met Leu Tyr Gly Gly
 1315 1320 1325

 agt gtt cct aat att act ttt gaa tcg gcg gca gaa gat tta agg cta 4032
 Ser Val Pro Asn Ile Thr Phe Glu Ser Ala Ala Glu Asp Leu Arg Leu
 1330 1335 1340

 tct acc aat atg gca ttg agt att att cat aat gga tat gcg gga acc 4080
 Ser Thr Asn Met Ala Leu Ser Ile Ile His Asn Gly Tyr Ala Gly Thr
 1345 1350 1355 1360

 cgc cgt ata caa tgt aat ctt atg aaa caa tac gct tca tta ggt gat 4128
 Arg Arg Ile Gln Cys Asn Leu Met Lys Gln Tyr Ala Ser Leu Gly Asp
 1365 1370 1375

 aaa ttt ata att tat gat tca tca ttt gat gat gca aac cgt ttt aat 4176
 Lys Phe Ile Ile Tyr Asp Ser Ser Phe Asp Asp Ala Asn Arg Phe Asn
 1380 1385 1390

 ctg gtg cca ttg ttt aaa ttc gga aaa gac gag aac tca gat gat agt 4224
 Leu Val Pro Leu Phe Lys Phe Gly Lys Asp Glu Asn Ser Asp Asp Ser
 1395 1400 1405

 att tgt ata tat aat gaa aac aac cct tcc tct gaa gat aag aag tgg tat 4272
 Ile Cys Ile Tyr Asn Glu Asn Pro Ser Ser Glu Asp Lys Lys Trp Tyr
 1410 1415 1420

 ttt tct tcg aaa gat gac aat aaa aca gcg gat tat aat ggt gga act 4320

Phe Ser Ser Lys Asp Asp Asn Lys Thr Ala Asp Tyr Asn Gly Gly Thr			
1425	1430	1435	1440
caa tgt ata gat gct gga acc agt aac aaa gat ttt tat tat aat ctc			4368
Gln Cys Ile Asp Ala Gly Thr Ser Asn Lys Asp Phe Tyr Tyr Asn Leu			
1445	1450	1455	
cag gag att gaa gta att agt gtt act ggt ggg tat tgg tcg agt tat			4416
Gln Glu Ile Glu Val Ile Ser Val Thr Gly Gly Tyr Trp Ser Ser Tyr			
1460	1465	1470	
aaa ata tcc aac ccg att aat atc aat acg ggc att gat agt gct aaa			4464
Lys Ile Ser Asn Pro Ile Asn Ile Asn Thr Gly Ile Asp Ser Ala Lys			
1475	1480	1485	
gta aaa gtc acc gta aaa gcg ggt ggt gac gat caa atc ttt act gct			4512
Val Lys Val Thr Val Lys Ala Gly Gly Asp Asp Gln Ile Phe Thr Ala			
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gat aat agt acc tat gtt cct cag caa ccg gca ccc agt ttt gag gag			4560
Asp Asn Ser Thr Tyr Val Pro Gln Gln Pro Ala Pro Ser Phe Glu Glu			
1505	1510	1515	1520
atg att tat cag ttc aat aac ctg aca ata gat tgt aag aat tta aat			4608
Met Ile Tyr Gln Phe Asn Asn Leu Thr Ile Asp Cys Lys Asn Leu Asn			
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ttc atc gac aat cag gca cat att gag att gat ttc acc gct acg gca			4656
Phe Ile Asp Asn Gln Ala His Ile Glu Ile Asp Phe Thr Ala Thr Ala			
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caa gat ggc cga ttc ttg ggt gca gaa act ttt att atc ccg gta act			4704
Gln Asp Gly Arg Phe Leu Gly Ala Glu Thr Phe Ile Ile Pro Val Thr			
1555	1560	1565	
aaa aaa gtt ctc ggt act gag aac gtg att gcg tta tat agc gaa aat			4752
Lys Lys Val Leu Gly Thr Glu Asn Val Ile Ala Leu Tyr Ser Glu Asn			
1570	1575	1580	
aac ggt gtt caa tat atg caa att ggc gca tat cgt acc cgt ttg aat			4800
Asn Gly Val Gln Tyr Met Gln Ile Gly Ala Tyr Arg Thr Arg Leu Asn			
1585	1590	1595	1600
acg tta ttc gct caa cag ttg gtt agc cgt gct aat cgt ggc att gat			4848
Thr Leu Phe Ala Gln Gln Leu Val Ser Arg Ala Asn Arg Gly Ile Asp			
1605	1610	1615	
gca gtg ctc agt atg gaa act cag aat att cag gaa ccg caa tta gga			4896
Ala Val Leu Ser Met Glu Thr Gln Asn Ile Gln Glu Pro Gln Leu Gly			
1620	1625	1630	
gcg ggc aca tat gtg cag ctt gtg ttg gat aaa tat gat gag tct att			4944
Ala Gly Thr Tyr Val Gln Leu Val Leu Asp Lys Tyr Asp Glu Ser Ile			
1635	1640	1645	
cat ggc act aat aaa agc ttt gct att gaa tat gtt gat ata ttt aaa			4992
His Gly Thr Asn Lys Ser Phe Ala Ile Glu Tyr Val Asp Ile Phe Lys			
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gag aac gat agt ttt gtg att tat caa gga gaa ctt agc gaa aca agt			5040
Glu Asn Asp Ser Phe Val Ile Tyr Gln Gly Glu Leu Ser Glu Thr Ser			

1665	1670	1675	1680	
caa act gtt gtg aaa gtt ttc tta tcc tat ttt ata gag gcg act gga Gln Thr Val Val Lys Val Phe Leu Ser Tyr Phe Ile Glu Ala Thr Gly				5088
1685		1690	1695	
aat aag aac cac tta tgg gta cgt gct aaa tac caa aag gaa acg act Asn Lys Asn His Leu Trp Val Arg Ala Lys Tyr Gln Lys Glu Thr Thr				5136
1700		1705	1710	
gat aag atc ttg ttc gac cgt act gat gag aaa gat ccg cac ggt tgg Asp Lys Ile Leu Phe Asp Arg Thr Asp Glu Lys Asp Pro His Gly Trp				5184
1715		1720	1725	
ttt ctc agc gac gat cac aag acc ttt agt ggt ctc tct tcc gca cag Phe Leu Ser Asp Asp His Lys Thr Phe Ser Gly Leu Ser Ser Ala Gln				5232
1730		1735	1740	
gca tta aag aac gac agt gaa ccg atg gat ttc tct ggc gcc aat gct Ala Leu Lys Asn Asp Ser Glu Pro Met Asp Phe Ser Gly Ala Asn Ala				5280
1745		1750	1755	1760
ctc tat ttc tgg gaa ctg ttc tat tac acg ccg atg atg atg gct cat Leu Tyr Phe Trp Glu Leu Phe Tyr Tyr Thr Pro Met Met Met Ala His				5328
1765		1770	1775	
cgt ttg ttg cag gaa cag aat ttt gat gcg gcg aac cat tgg ttc cgt Arg Leu Leu Gln Glu Gln Asn Phe Asp Ala Ala Asn His Trp Phe Arg				5376
1780		1785	1790	
tat gtc tgg agt cca tcc ggt tat atc gtt gat ggt aaa att gct atc Tyr Val Trp Ser Pro Ser Gly Tyr Ile Val Asp Gly Lys Ile Ala Ile				5424
1795		1800	1805	
tac cac tgg aac gtg cga ccg ctg gaa gaa gac acc agt tgg aat gca Tyr His Trp Asn Val Arg Pro Leu Glu Asp Thr Ser Trp Asn Ala				5472
1810		1815	1820	
caa caa ctg gac tcc acc gat cca gat gct gta gcc caa gat gat ccg Gln Gln Leu Asp Ser Thr Asp Pro Asp Ala Val Ala Gln Asp Asp Pro				5520
1825		1830	1835	1840
atg cac tac aag gtg gct acc ttt atg gcg acg ttg gat ctg cta atg Met His Tyr Lys Val Ala Thr Phe Met Ala Thr Leu Asp Leu Met				5568
1845		1850	1855	
gcc cgt ggt gat gct gct tac cgc cag tta gag cgt gat acg ttg gct Ala Arg Gly Asp Ala Ala Tyr Arg Gln Leu Glu Arg Asp Thr Leu Ala				5616
1860		1865	1870	
gaa gct aaa atg tgg tat aca cag gcg ctt aat ctg ttg ggt gat gag Glu Ala Lys Met Trp Tyr Thr Gln Ala Leu Asn Leu Leu Gly Asp Glu				5664
1875		1880	1885	
cca caa gtg atg ctg agt acg act tgg gct aat cca aca ttg ggt aat Pro Gln Val Met Leu Ser Thr Thr Trp Ala Asn Pro Thr Leu Gly Asn				5712
1890		1895	1900	
gct gct tca aaa acc aca cag cag gtt cgt cag caa gtg ctt acc cag Ala Ala Ser Lys Thr Thr Gln Gln Val Arg Gln Gln Val Leu Thr Gln				5760
1905		1910	1915	1920

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Leu Arg Leu Asn Ser Arg Val Lys Thr Pro Leu Leu Gly Thr Ala Asn	
1925 1930 1935	
tcc ctg acc gct tta ttc ctg ccg cag gaa aat agc aag ctc aaa ggc	5856
Ser Leu Thr Ala Leu Phe Leu Pro Gln Glu Asn Ser Lys Leu Lys Gly	
1940 1945 1950	
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Tyr Trp Arg Thr Leu Ala Gln Arg Met Phe Asn Leu Arg His Asn Leu	
1955 1960 1965	
tcg att gac ggc cag ccg ctc tcc ttg ccg ctg tat gct aaa ccg gct	5952
Ser Ile Asp Gly Gln Pro Leu Ser Leu Pro Leu Tyr Ala Lys Pro Ala	
1970 1975 1980	
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Asp Pro Lys Ala Leu Leu Ser Ala Ala Val Ser Ala Ser Gln Gly Gly	
1985 1990 1995 2000	
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Ala Asp Leu Pro Lys Ala Pro Leu Thr Ile His Arg Phe Pro Gln Met	
2005 2010 2015	
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Leu Glu Gly Ala Arg Gly Leu Val Asn Gln Leu Ile Gln Phe Gly Ser	
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tca cta ttg ggg tac agt gag cgt cag gat gcg gaa gct atg agt caa	6144
Ser Leu Leu Gly Tyr Ser Glu Arg Gln Asp Ala Glu Ala Met Ser Gln	
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Leu Leu Gln Thr Gln Ala Ser Glu Leu Ile Leu Thr Ser Ile Arg Met	
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cag gat aac caa ttg gca gag ctg gat tcg gaa aaa acc gcc ttg caa	6240
Gln Asp Asn Gln Leu Ala Glu Leu Asp Ser Glu Lys Thr Ala Leu Gln	
2065 2070 2075 2080	
gtc tct tta gct gga gtg caa caa ccg ttt gac agc tat agc caa ctg	6288
Val Ser Leu Ala Gly Val Gln Gln Arg Phe Asp Ser Tyr Ser Gln Leu	
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tat gag gag aac atc aac gca ggt gag cag cga gcg ctg gcg ttg cgc	6336
Tyr Glu Glu Asn Ile Asn Ala Gly Glu Gln Arg Ala Leu Ala Leu Arg	
2100 2105 2110	
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Ser Glu Ser Ala Ile Glu Ser Gln Gly Ala Gln Ile Ser Arg Met Ala	
2115 2120 2125	
ggc gcg ggt gtt gat atg gca cca aat atc ttc ggc ctg gct gat ggc	6432
Gly Ala Gly Val Asp Met Ala Pro Asn Ile Phe Gly Leu Ala Asp Gly	
2130 2135 2140	
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Gly Met His Tyr Gly Ala Ile Ala Tyr Ala Ile Ala Asp Gly Ile Glu	
2145 2150 2155 2160	

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 Leu Ser Ala Ser Ala Lys Met Val Asp Ala Glu Lys Val Ala Gln Ser
 2165 2170 2175

gaa ata tat cgc cgt cgc cgt caa gaa tgg aaa att cag cgt gac aac 6576
 Glu Ile Tyr Arg Arg Arg Gln Glu Trp Lys Ile Gln Arg Asp Asn
 2180 2185 2190

gca caa gcg gag att aac cag tta aac gcg caa ctg gaa tca ctg tct 6624
 Ala Gln Ala Glu Ile Asn Gln Leu Asn Ala Gln Leu Glu Ser Leu Ser
 2195 2200 2205

att cgc cgt gaa gcc gct gaa atg caa aaa gag tac ctg aaa acc cag 6672
 Ile Arg Arg Glu Ala Ala Glu Met Gln Lys Glu Tyr Leu Lys Thr Gln
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aat caa gcg tta tat agt tgg tta cga ggg cgt ttg tca ggt att tat 6768
 Asn Gln Ala Leu Tyr Ser Trp Leu Arg Gly Arg Leu Ser Gly Ile Tyr
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 Phe Gln Phe Tyr Asp Leu Ala Val Ser Arg Cys Leu Met Ala Glu Gln
 2260 2265 2270

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 Ser Tyr Gln Trp Glu Ala Asn Asp Asn Ser Ile Ser Phe Val Lys Pro
 2275 2280 2285

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 Arg Ala Leu Glu Val Glu Arg Thr Val Ser Leu Ala Val Val Tyr Asp
 2325 2330 2335

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 Ser Leu Glu Gly Asn Asp Arg Phe Asn Leu Ala Glu Gln Ile Pro Ala
 2340 2345 2350

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 Leu Leu Asp Lys Gly Glu Gly Thr Ala Gly Thr Lys Glu Asn Gly Leu
 2355 2360 2365

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 Ser Leu Ala Asn Ala Ile Leu Ser Ala Ser Val Lys Leu Ser Asp Leu
 2370 2375 2380

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 Lys Leu Gly Thr Asp Tyr Pro Asp Ser Ile Val Gly Ser Asn Lys Val
 2385 2390 2395 2400

cgt cgt att aag caa atc agt gtt tcg cta cct gca ttg gtt ggg cct 7248

Arg Arg Ile Lys Gln Ile Ser Val Ser Leu Pro Ala Leu Val Gly Pro		
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tat cag gat gtt cag gct atg ctc agc tat ggt ggc agt act caa ttg		7296
Tyr Gln Asp Val Gln Ala Met Leu Ser Tyr Gly Gly Ser Thr Gln Leu		
2420	2425	2430
ccg aaa ggt tgt tca gcg ttg gct gtg tct cat ggt acc aat gat agt		7344
Pro Lys Gly Cys Ser Ala Leu Ala Val Ser His Gly Thr Asn Asp Ser		
2435	2440	2445
ggc cag ttc cag ttg gat ttc aat gac ggc aaa tac ctg cca ttt gaa		7392
Gly Gln Phe Gln Leu Asp Phe Asn Asp Gly Lys Tyr Leu Pro Phe Glu		
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ggc att gct ctt gat gat cag ggt aca ctg aat ctt caa ttt ccg aat		7440
Gly Ile Ala Leu Asp Asp Gln Gly Thr Leu Asn Leu Gln Phe Pro Asn		
2465	2470	2475
gct acc gac aag cag aaa gca ata ttg caa act atg agc gat att att		7488
Ala Thr Asp Lys Gln Lys Ala Ile Leu Gln Thr Met Ser Asp Ile Ile		
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Gln Cys Gly Phe Asn Cys Leu Thr Asp Ile Ser His Ser Ser Phe Asn		
20	25	30
gag ttc aga caa caa gtc tct gag cac ctc tcc tgg tcc gag acc cat		143
Glu Phe Arg Gln Gln Val Ser Glu His Leu Ser Trp Ser Glu Thr His		
35	40	45
gac ctc tac cat gac gct cag caa gct cag aag gac aac agg ctc tac		191
Asp Leu Tyr His Asp Ala Gln Gln Ala Gln Lys Asp Asn Arg Leu Tyr		
50	55	60
gag gct agg atc ctc aag agg gct aac cca caa ctc cag aac gct gtc		239
Glu Ala Arg Ile Leu Lys Arg Ala Asn Pro Gln Leu Gln Asn Ala Val		
65	70	75

cac ctc gcc atc ttg gct cca aac gct gag ttg att ggt tac aac aac	80	85	90	95	287
His Leu Ala Ile Leu Ala Pro Asn Ala Glu Leu Ile Gly Tyr Asn Asn					
cag ttc tct ggc aga gct agc cag tac gtg gct cct ggt aca gtc tcc	100	105	110		335
Gln Phe Ser Gly Arg Ala Ser Gln Tyr Val Ala Pro Gly Thr Val Ser					
tcc atg ttc agc cca gcc gct tac ctc act gag ttg tac cgc gag gct	115	120	125		383
Ser Met Phe Ser Pro Ala Ala Tyr Leu Thr Glu Leu Tyr Arg Glu Ala					
agg aac ctt cat gct tct gac tcc gtc tac tac ttg gac aca cgc aga	130	135	140		431
Arg Asn Leu His Ala Ser Asp Ser Val Tyr Tyr Leu Asp Thr Arg Arg					
cca gac ctc aag agc atg gcc ctc agc caa cag aac atg gac att gag	145	150	155		479
Pro Asp Leu Lys Ser Met Ala Leu Ser Gln Gln Asn Met Asp Ile Glu					
ttg tcc acc ctc tcc ttg agc aac gag ctt ctc ttg gag tcc atc aag	160	165	170	175	527
Leu Ser Thr Leu Ser Leu Ser Asn Glu Leu Leu Leu Glu Ser Ile Lys					
act gag agc aag ttg gag aac tac acc aag gtc atg gag atg ctc tcc	180	185	190		575
Thr Glu Ser Lys Leu Glu Asn Tyr Thr Lys Val Met Glu Met Leu Ser					
acc ttc aga cca agc ggt gca act cca tac cat gat gcc tac gag aac	195	200	205		623
Thr Phe Arg Pro Ser Gly Ala Thr Pro Tyr His Asp Ala Tyr Glu Asn					
gtc agg gag gtc atc caa ctt caa gac cct ggt ctt gag caa ctc aac	210	215	220		671
Val Arg Glu Val Ile Gln Leu Gln Asp Pro Gly Leu Glu Gln Leu Asn					
gct tct cca gcc att gct ggt ttg atg cac cag gca tcc ttg ctc ggt	225	230	235		719
Ala Ser Pro Ala Ile Ala Gly Leu Met His Gln Ala Ser Leu Leu Gly					
atc aac gcc tcc atc tct cct gag ttg ttc aac atc ttg act gag gag	240	245	250	255	767
Ile Asn Ala Ser Ile Ser Pro Glu Leu Phe Asn Ile Leu Thr Glu Glu					
atc act gag ggc aac gct gag gag ttg tac aag aag aac ttc ggc aac	260	265	270		815
Ile Thr Glu Gly Asn Ala Glu Glu Leu Tyr Lys Lys Asn Phe Gly Asn					
att gag cca gcc tct ctt gca atg cct gag tac ctc aag agg tac tac	275	280	285		863
Ile Glu Pro Ala Ser Leu Ala Met Pro Glu Tyr Leu Lys Arg Tyr Tyr					
aac ttg tct gat gag gag ctt tct caa ttc att ggc aag gct tcc aac	290	295	300		911
Asn Leu Ser Asp Glu Glu Leu Ser Gln Phe Ile Gly Lys Ala Ser Asn					
ttc ggt caa cag gag tac agc aac aac cag ctc atc act cca gtt gtg	305	310	315		959
Phe Gly Gln Gln Glu Tyr Ser Asn Asn Gln Leu Ile Thr Pro Val Val					

aac tcc tct gat ggc act gtg aag gtc tac cgc atc aca cgt gag tac 1007
 Asn Ser Ser Asp Gly Thr Val Lys Val Tyr Arg Ile Thr Arg Glu Tyr
 320 325 330 335

 acc aca aac gcc tac caa atg gat gtt gag ttg ttc cca ttc ggt ggt 1055
 Thr Thr Asn Ala Tyr Gln Met Asp Val Glu Leu Phe Pro Phe Gly Gly
 340 345 350

 gag aac tac aga ctt gac tac aag ttc aag aac ttc tac aac gcc tcc 1103
 Glu Asn Tyr Arg Leu Asp Tyr Lys Phe Lys Asn Phe Tyr Asn Ala Ser
 355 360 365

 tac ctc tcc atc aag ttg aac gac aag agg gag ctt gtc agg act gag 1151
 Tyr Leu Ser Ile Lys Leu Asn Asp Lys Arg Glu Leu Val Arg Thr Glu
 370 375 380

 ggt gct cct caa gtg aac att gag tac tct gcc aac atc acc ctc aac 1199
 Gly Ala Pro Gln Val Asn Ile Glu Tyr Ser Ala Asn Ile Thr Leu Asn
 385 390 395

 aca gct gac atc tct caa cca ttc gag att ggt ttg acc aga gtc ctt 1247
 Thr Ala Asp Ile Ser Gln Pro Phe Glu Ile Gly Leu Thr Arg Val Leu
 400 405 410 415

 ccc tct ggc tcc tgg gcc tac gct gca gcc aag ttc act gtt gag gag 1295
 Pro Ser Gly Ser Trp Ala Tyr Ala Ala Lys Phe Thr Val Glu Glu
 420 425 430

 tac aac cag tac tct ttc ctc ttg aag ctc aac aag gca att cgt ctc 1343
 Tyr Asn Gln Tyr Ser Phe Leu Leu Lys Leu Asn Lys Ala Ile Arg Leu
 435 440 445

 agc aga gcc act gag ttg tct ccc acc atc ttg gag ggc att gtg agg 1391
 Ser Arg Ala Thr Glu Leu Ser Pro Thr Ile Leu Glu Gly Ile Val Arg
 450 455 460

 tct gtc aac ctt caa ctt gac atc aac act gat gtg ctt ggc aag gtc 1439
 Ser Val Asn Leu Gln Leu Asp Ile Asn Thr Asp Val Leu Gly Lys Val
 465 470 475

 ttc ctc acc aag tac tac atg caa cgc tac gcc atc cat gct gag act 1487
 Phe Leu Thr Lys Tyr Tyr Met Gln Arg Tyr Ala Ile His Ala Glu Thr
 480 485 490 495

 gca ctc atc ctc tgc aac gca ccc atc tct caa cgc tcc tac gac aac 1535
 Ala Leu Ile Leu Cys Asn Ala Pro Ile Ser Gln Arg Ser Tyr Asp Asn
 500 505 510

 cag cct tcc cag ttc gac agg ctc ttc aac act cct ctc ttg aac ggc 1583
 Gln Pro Ser Gln Phe Asp Arg Leu Phe Asn Thr Pro Leu Leu Asn Gly
 515 520 525

 cag tac ttc tcc act ggt gat gag gag att gac ctc aac tct ggc tcc 1631
 Gln Tyr Phe Ser Thr Gly Asp Glu Glu Ile Asp Leu Asn Ser Gly Ser
 530 535 540

 aca ggt gac tgg aga aag acc atc ttg aag agg gcc ttc aac att gat 1679
 Thr Gly Asp Trp Arg Lys Thr Ile Leu Lys Arg Ala Phe Asn Ile Asp
 545 550 555

 gat gtc tct ctc ttc cgt ctc ttg aag atc aca gat cac gac aac aag 1727

Asp Val Ser Leu Phe Arg Leu Leu Lys Ile Thr Asp His Asp Asn Lys				
560	565	570	575	
gat ggc aag atc aag aac aac ttg aag aac ctt tcc aac ctc tac att				1775
Asp Gly Lys Ile Lys Asn Asn Leu Lys Asn Leu Ser Asn Leu Tyr Ile				
580	585	590		
ggc aag ttg ctt gca gac atc cac caa ctc acc att gat gag ttg gac				1823
Gly Lys Leu Leu Ala Asp Ile His Gln Leu Thr Ile Asp Glu Leu Asp				
595	600	605		
ctc ttg ctc att gca gtc ggt gag ggc aag acc aac ctc tct gca atc				1871
Leu Leu Leu Ile Ala Val Gly Glu Gly Lys Thr Asn Leu Ser Ala Ile				
610	615	620		
tct gac aag cag ttg gca acc ctc atc agg aag ttg aac acc atc acc				1919
Ser Asp Lys Gln Leu Ala Thr Leu Ile Arg Lys Leu Asn Thr Ile Thr				
625	630	635		
tcc tgg ctt cac acc cag aag tgg tct gtc ttc caa ctc ttc atc atg				1967
Ser Trp Leu His Thr Gln Lys Trp Ser Val Phe Gln Leu Phe Ile Met				
640	645	650	655	
acc agc acc tcc tac aac aag acc ctc act cct gag atc aag aac ctc				2015
Thr Ser Thr Ser Tyr Asn Lys Thr Leu Thr Pro Glu Ile Lys Asn Leu				
660	665	670		
ttg gac aca gtc tac cac ggt ctc caa ggc ttc gac aag gac aag gct				2063
Leu Asp Thr Val Tyr His Gly Leu Gln Gly Phe Asp Lys Asp Lys Ala				
675	680	685		
gac ttg ctt cat gtc atg gct ccc tac att gca gcc acc ctc caa ctc				2111
Asp Leu Leu His Val Met Ala Pro Tyr Ile Ala Ala Thr Leu Gln Leu				
690	695	700		
tcc tct gag aac gtg gct cac tct gtc ttg ctc tgg gct gac aag ctc				2159
Ser Ser Glu Asn Val Ala His Ser Val Leu Leu Trp Ala Asp Lys Leu				
705	710	715		
caa cct ggt gat ggt gcc atg act gct gag aag ttc tgg gac tgg ctc				2207
Gln Pro Gly Asp Gly Ala Met Thr Ala Glu Lys Phe Trp Asp Trp Leu				
720	725	730	735	
aac acc aag tac aca cca ggc tcc tct gag gct gtt gag act caa gag				2255
Asn Thr Lys Tyr Thr Pro Gly Ser Ser Glu Ala Val Glu Thr Gln Glu				
740	745	750		
cac att gtg caa tac tgc cag gct ctt gca cag ttg gag atg gtc tac				2303
His Ile Val Gln Tyr Cys Gln Ala Leu Ala Gln Leu Glu Met Val Tyr				
755	760	765		
cac tcc act ggc atc aac gag aac gct ttc aga ctc ttc gtc acc aag				2351
His Ser Thr Gly Ile Asn Glu Asn Ala Phe Arg Leu Phe Val Thr Lys				
770	775	780		
cct gag atg ttc ggt gct gcc aca ggt gct gca cct gct cat gat gct				2399
Pro Glu Met Phe Gly Ala Ala Thr Gly Ala Ala Pro Ala His Asp Ala				
785	790	795		
ctc tcc ctc atc atg ttg acc agg ttc gct gac tgg gtc aac gct ctt				2447
Leu Ser Leu Ile Met Leu Thr Arg Phe Ala Asp Trp Val Asn Ala Leu				

800	805	810	815	
ggt gag aag gct tcc tct gtc ttg gct gcc ttc gag gcc aac tcc ctc				2495
Gly Glu Lys Ala Ser Ser Val Leu Ala Ala Phe Glu Ala Asn Ser Leu				
820	825	830		
act gct gag caa ctt gct gat gcc atg aac ctt gat gcc aac ctc ttg				2543
Thr Ala Glu Gln Leu Ala Asp Ala Met Asn Leu Asp Ala Asn Leu Leu				
835	840	845		
ctc caa gct tcc att caa gct cag aac cac caa cac ctc cca cct gtc				2591
Leu Gln Ala Ser Ile Gln Ala Gln Asn His Gln His Leu Pro Pro Val				
850	855	860		
act cca gag aac gct ttc tcc tgc tgg acc tcc atc aac acc atc ctc				2639
Thr Pro Glu Asn Ala Phe Ser Cys Trp Thr Ser Ile Asn Thr Ile Leu				
865	870	875		
caa tgg gtc aac gtg gct cag caa ctc aac gtg gct cca caa ggt gtc				2687
Gln Trp Val Asn Val Ala Gln Gln Leu Asn Val Ala Pro Gln Gly Val				
880	885	890	895	
tct gct ttg gtc ggt ctt gac tac atc cag tcc atg aag gag aca cca				2735
Ser Ala Leu Val Gly Leu Asp Tyr Ile Gln Ser Met Lys Glu Thr Pro				
900	905	910		
acc tac gct caa tgg gag aac gca gct ggt gtc ttg act gct ggt ctc				2783
Thr Tyr Ala Gln Trp Glu Asn Ala Ala Gly Val Leu Thr Ala Gly Leu				
915	920	925		
aac tcc caa cag gcc aac acc ctc cat gct ttc ttg gat gag tct cgc				2831
Asn Ser Gln Gln Ala Asn Thr Leu His Ala Phe Leu Asp Glu Ser Arg				
930	935	940		
tct gct gcc ctc tcc acc tac tac atc agg caa gtc gcc aag gca gct				2879
Ser Ala Ala Leu Ser Thr Tyr Tyr Ile Arg Gln Val Ala Lys Ala Ala				
945	950	955		
gct gcc atc aag tct cgc gat gac ctc tac caa tac ctc ctc att gac				2927
Ala Ala Ile Lys Ser Arg Asp Asp Leu Tyr Gln Tyr Leu Leu Ile Asp				
960	965	970	975	
aac cag gtc tct gct gcc atc aag acc acc agg atc gct gag gcc atc				2975
Asn Gln Val Ser Ala Ala Ile Lys Thr Thr Arg Ile Ala Glu Ala Ile				
980	985	990		
gct tcc atc caa ctc tac gtc aac cgc gct ctt gag aac gtt gag gag				3023
Ala Ser Ile Gln Leu Tyr Val Asn Arg Ala Leu Glu Asn Val Glu Glu				
995	1000	1005		
aac gcc aac tct ggt gtc atc tct cgc caa ttc ttc atc gac tgg gac				3071
Asn Ala Asn Ser Gly Val Ile Ser Arg Gln Phe Phe Ile Asp Trp Asp				
1010	1015	1020		
aag tac aac aag agg tac tcc acc tgg gct ggt gtc tct caa ctt gtc				3119
Lys Tyr Asn Lys Arg Tyr Ser Thr Trp Ala Gly Val Ser Gln Leu Val				
1025	1030	1035		
tac tac cca gag aac tac att gac cca acc atg agg att ggt cag acc				3167
Tyr Tyr Pro Glu Asn Tyr Ile Asp Pro Thr Met Arg Ile Gly Gln Thr				
1040	1045	1050	1055	

aag atg atg gat gct ctc ttg caa tct gtc tcc caa agc caa ctc aac Lys Met Met Asp Ala Leu Leu Gln Ser Val Ser Gln Ser Gln Leu Asn 1060 1065 1070	3215
gct gac act gtg gag gat gcc ttc atg agc tac ctc acc tcc ttc gag Ala Asp Thr Val Glu Asp Ala Phe Met Ser Tyr Leu Thr Ser Phe Glu 1075 1080 1085	3263
caa gtt gcc aac ctc aag gtc atc tct gct tac cat gac aac atc aac Gln Val Ala Asn Leu Lys Val Ile Ser Ala Tyr His Asp Asn Ile Asn 1090 1095 1100	3311
aac gac caa ggt ctc acc tac ttc att ggt ctc tct gag act gat gct Asn Asp Gln Gly Leu Thr Tyr Phe Ile Gly Leu Ser Glu Thr Asp Ala 1105 1110 1115	3359
ggt gag tac tac tgg aga tcc gtg gac cac agc aag ttc aac gat ggc Gly Glu Tyr Tyr Trp Arg Ser Val Asp His Ser Lys Phe Asn Asp Gly 1120 1125 1130 1135	3407
aag ttc gct gca aac gct tgg tct gag tgg cac aag att gac tgc cct Lys Phe Ala Ala Asn Ala Trp Ser Glu Trp His Lys Ile Asp Cys Pro 1140 1145 1150	3455
atc aac cca tac aag tcc acc atc aga cct gtc atc tac aag agc cgc Ile Asn Pro Tyr Lys Ser Thr Ile Arg Pro Val Ile Tyr Lys Ser Arg 1155 1160 1165	3503
ctc tac ttg ctc tgg ctt gag cag aag gag atc acc aag caa act ggc Leu Tyr Leu Leu Trp Leu Glu Gln Lys Glu Ile Thr Lys Gln Thr Gly 1170 1175 1180	3551
aac tcc aag gat ggt tac caa act gag act gac tac cgc tac gag ttg Asn Ser Lys Asp Gly Tyr Gln Thr Glu Thr Asp Tyr Arg Tyr Glu Leu 1185 1190 1195	3599
aag ttg gct cac atc cgc tac gat ggt acc tgg aac act cca atc acc Lys Leu Ala His Ile Arg Tyr Asp Gly Thr Trp Asn Thr Pro Ile Thr 1200 1205 1210 1215	3647
ttc gat gtc aac aag aag atc agc gag ttg aag ttg gag aag aac cgt Phe Asp Val Asn Lys Lys Ile Ser Glu Leu Lys Leu Glu Lys Asn Arg 1220 1225 1230	3695
gct cct ggt ctc tac tgc gct ggt tac caa ggt gag gac acc ctc ttg Ala Pro Gly Leu Tyr Cys Ala Gly Tyr Gln Gly Glu Asp Thr Leu Leu 1235 1240 1245	3743
gtc atg ttc tac aac cag caa gac acc ctt gac tcc tac aag aac gct Val Met Phe Tyr Asn Gln Gln Asp Thr Leu Asp Ser Tyr Lys Asn Ala 1250 1255 1260	3791
tcc atg caa ggt ctc tac atc ttc gct gac atg gct tcc aag gac atg Ser Met Gln Gly Leu Tyr Ile Phe Ala Asp Met Ala Ser Lys Asp Met 1265 1270 1275	3839
act cca gag caa agc aac gtc tac cgt gac aac tcc tac caa cag ttc Thr Pro Glu Gln Ser Asn Val Tyr Arg Asp Asn Ser Tyr Gln Gln Phe 1280 1285 1290 1295	3887

gac acc aac aac gtc agg cgt gtc aac aac aga tac gct gag gac tac 3935
 Asp Thr Asn Asn Val Arg Arg Val Asn Asn Arg Tyr Ala Glu Asp Tyr
 1300 1305 1310

gag atc cca agc tct gtc agc tct cgc aag gac tac ggc tgg ggt gac 3983
 Glu Ile Pro Ser Ser Val Ser Arg Lys Asp Tyr Gly Trp Gly Asp
 1315 1320 1325

tac tac ctc agc atg gtg tac aac ggt gac atc cca acc atc aac tac 4031
 Tyr Tyr Leu Ser Met Val Tyr Asn Gly Asp Ile Pro Thr Ile Asn Tyr
 1330 1335 1340

aag gct gcc tct tcc gac ctc aaa atc tac atc agc cca aag ctc agg 4079
 Lys Ala Ala Ser Ser Asp Leu Lys Ile Tyr Ile Ser Pro Lys Leu Arg
 1345 1350 1355

atc atc cac aac ggc tac gag ggt cag aag agg aac cag tgc aac ttg 4127
 Ile Ile His Asn Gly Tyr Glu Gly Gln Lys Arg Asn Gln Cys Asn Leu
 1360 1365 1370 1375

atg aac aag tac ggc aag ttg ggt gac aag ttc att gtc tac acc tct 4175
 Met Asn Lys Tyr Gly Lys Leu Gly Asp Lys Phe Ile Val Tyr Thr Ser
 1380 1385 1390

ctt ggt gtc aac cca aac aac agc tcc aac aag ctc atg ttc tac cca 4223
 Leu Gly Val Asn Pro Asn Asn Ser Ser Asn Lys Leu Met Phe Tyr Pro
 1395 1400 1405

gtc tac caa tac tct ggc aac acc tct ggt ctc aac cag ggt aga ctc 4271
 Val Tyr Gln Tyr Ser Gly Asn Thr Ser Gly Leu Asn Gln Gly Arg Leu
 1410 1415 1420

ttg ttc cac agg gac acc acc tac cca agc aag gtg gag gct tgg att 4319
 Leu Phe His Arg Asp Thr Thr Tyr Pro Ser Lys Val Glu Ala Trp Ile
 1425 1430 1435

cct ggt gcc aag agg tcc ctc acc aac cag aac gct gcc att ggt gat 4367
 Pro Gly Ala Lys Arg Ser Leu Thr Asn Gln Asn Ala Ala Ile Gly Asp
 1440 1445 1450 1455

gac tac gcc aca gac tcc ctc aac aag cct gat gac ctc aag cag tac 4415
 Asp Tyr Ala Thr Asp Ser Leu Asn Lys Pro Asp Asp Leu Lys Gln Tyr
 1460 1465 1470

atc ttc atg act gac tcc aag ggc aca gcc act gat gtc tct ggt cca 4463
 Ile Phe Met Thr Asp Ser Lys Gly Thr Ala Thr Asp Val Ser Gly Pro
 1475 1480 1485

gtg gag atc aac act gca atc agc cca gcc aag gtc caa atc att gtc 4511
 Val Glu Ile Asn Thr Ala Ile Ser Pro Ala Lys Val Gln Ile Ile Val
 1490 1495 1500

aag gct ggt ggc aag gag caa acc ttc aca gct gac aag gat gtc tcc 4559
 Lys Ala Gly Gly Lys Glu Gln Thr Phe Thr Ala Asp Lys Asp Val Ser
 1505 1510 1515

atc cag cca agc cca tcc ttc gat gag atg aac tac caa ttc aac gct 4607
 Ile Gln Pro Ser Pro Ser Phe Asp Glu Met Asn Tyr Gln Phe Asn Ala
 1520 1525 1530 1535

ctt gag att gat ggt tct ggc ctc aac ttc atc aac aac tct gct tcc 4655

Leu Glu Ile Asp Gly Ser Gly Leu Asn Phe Ile Asn Asn Ser Ala Ser			
1540	1545	1550	
att gat gtc acc ttc act gcc ttc gct gag gat ggc cgc aag ttg ggt			4703
Ile Asp Val Thr Phe Thr Ala Phe Ala Glu Asp Gly Arg Lys Leu Gly			
1555	1560	1565	
tac gag agc ttc tcc atc cca gtc acc ctt aag gtt tcc act gac aac			4751
Tyr Glu Ser Phe Ser Ile Pro Val Thr Leu Lys Val Ser Thr Asp Asn			
1570	1575	1580	
gca ctc acc ctt cat cac aac gag aac ggt gct cag tac atg caa tgg			4799
Ala Leu Thr Leu His His Asn Glu Asn Gly Ala Gln Tyr Met Gln Trp			
1585	1590	1595	
caa agc tac cgc acc agg ttg aac acc ctc ttc gca agg caa ctt gtg			4847
Gln Ser Tyr Arg Thr Arg Leu Asn Thr Leu Phe Ala Arg Gln Leu Val			
1600	1605	1610	1615
gcc cgt gcc acc aca ggc att gac acc atc ctc agc atg gag acc cag			4895
Ala Arg Ala Thr Thr Gly Ile Asp Thr Ile Leu Ser Met Glu Thr Gln			
1620	1625	1630	
aac atc caa gag cca cag ttg ggc aag ggt ttc tac gcc acc ttc gtc			4943
Asn Ile Gln Glu Pro Gln Leu Gly Lys Gly Phe Tyr Ala Thr Phe Val			
1635	1640	1645	
atc cca cct tac aac ctc agc act cat ggt gat gag agg tgg ttc aag			4991
Ile Pro Pro Tyr Asn Leu Ser Thr His Gly Asp Glu Arg Trp Phe Lys			
1650	1655	1660	
ctc tac atc aag cac gtg gtt gac aac aac tcc cac atc atc tac tct			5039
Leu Tyr Ile Lys His Val Val Asp Asn Asn Ser His Ile Ile Tyr Ser			
1665	1670	1675	
ggt caa ctc act gac acc aac atc aac atc acc ctc ttc atc cca ctt			5087
Gly Gln Leu Thr Asp Thr Asn Ile Asn Ile Thr Leu Phe Ile Pro Leu			
1680	1685	1690	1695
gac gat gtc cca ctc aac cag gac tac cat gcc aag gtc tac atg acc			5135
Asp Asp Val Pro Leu Asn Gln Asp Tyr His Ala Lys Val Tyr Met Thr			
1700	1705	1710	
ttc aag aag tct cca tct gat ggc acc tgg tgg ggt cca cac ttc gtc			5183
Phe Lys Lys Ser Pro Ser Asp Gly Thr Trp Trp Gly Pro His Phe Val			
1715	1720	1725	
cgt gat gac aag ggc atc gtc acc atc aac cca aag tcc atc ctc acc			5231
Arg Asp Asp Lys Gly Ile Val Thr Ile Asn Pro Lys Ser Ile Leu Thr			
1730	1735	1740	
cac ttc gag tct gtc aac gtt ctc aac aac atc tcc tct gag cca atg			5279
His Phe Glu Ser Val Asn Val Leu Asn Asn Ile Ser Ser Glu Pro Met			
1745	1750	1755	
gac ttc tct ggt gcc aac tcc ctc tac ttc tgg gag ttg ttc tac tac			5327
Asp Phe Ser Gly Ala Asn Ser Leu Tyr Phe Trp Glu Leu Phe Tyr Tyr			
1760	1765	1770	1775
aca cca atg ctt gtg gct caa agg ttg ctc cat gag cag aac ttc gat			5375
Thr Pro Met Leu Val Ala Gln Arg Leu Leu His Glu Gln Asn Phe Asp			

1780	1785	1790	
gag gcc aac agg tgg ctc aag tac gtc tgg agc cca tct ggt tac att Glu Ala Asn Arg Trp Leu Lys Tyr Val Trp Ser Pro Ser Gly Tyr Ile 1795 1800 1805			5423
gtg cat ggt caa atc cag aac tac caa tgg aac gtc agg cca ttg ctt Val His Gly Gln Ile Gln Asn Tyr Gln Trp Asn Val Arg Pro Leu Leu 1810 1815 1820			5471
gag gac acc tcc tgg aac tct gac cca ctt gac tct gtg gac cct gat Glu Asp Thr Ser Trp Asn Ser Asp Pro Leu Asp Ser Val Asp Pro Asp 1825 1830 1835			5519
gct gtg gct caa cat gac cca atg cac tac aag gtc tcc acc ttc atg Ala Val Ala Gln His Asp Pro Met His Tyr Lys Val Ser Thr Phe Met 1840 1845 1850 1855			5567
agg acc ttg gac ctc ttg att gcc aga ggt gac cat gct tac cgc caa Arg Thr Leu Asp Leu Leu Ile Ala Arg Gly Asp His Ala Tyr Arg Gln 1860 1865 1870			5615
ttg gag agg gac acc ctc aac gag gca aag atg tgg tac atg caa gct Leu Glu Arg Asp Thr Leu Asn Glu Ala Lys Met Trp Tyr Met Gln Ala 1875 1880 1885			5663
ctc cac ctc ttg ggt gac aag cca tac ctc cca ctc agc acc act tgg Leu His Leu Leu Gly Asp Lys Pro Tyr Leu Pro Leu Ser Thr Thr Trp 1890 1895 1900			5711
tcc gac cca agg ttg gac cgt gct gac atc acc act cag aac gct Ser Asp Pro Arg Leu Asp Arg Ala Ala Asp Ile Thr Thr Gln Asn Ala 1905 1910 1915			5759
cat gac tct gcc att gtt gct ctc agg cag aac atc cca act cct gct His Asp Ser Ala Ile Val Ala Leu Arg Gln Asn Ile Pro Thr Pro Ala 1920 1925 1930 1935			5807
cca ctc tcc ctc aga tct gct aac acc ctc act gac ttg ttc ctc cca Pro Leu Ser Leu Arg Ser Ala Asn Thr Leu Thr Asp Leu Phe Leu Pro 1940 1945 1950			5855
cag atc aac gag gtc atg atg aac tac tgg caa acc ttg gct caa agg Gln Ile Asn Glu Val Met Met Asn Tyr Trp Gln Thr Leu Ala Gln Arg 1955 1960 1965			5903
gtc tac aac ctc aga cac aac ctc tcc att gat ggt caa cca ctc tac Val Tyr Asn Leu Arg His Asn Leu Ser Ile Asp Gly Gln Pro Leu Tyr 1970 1975 1980			5951
ctc cca atc tac gcc aca cca gct gac cca aag gct ctt ctc tct gct Leu Pro Ile Tyr Ala Thr Pro Ala Asp Pro Lys Ala Leu Leu Ser Ala 1985 1990 1995			5999
gct gtg gct acc agc caa ggt ggt ggc aag ctc cca gag tcc ttc atg Ala Val Ala Thr Ser Gln Gly Gly Gly Lys Leu Pro Glu Ser Phe Met 2000 2005 2010 2015			6047
tcc ctc tgg agg ttc cca cac atg ttg gag aac gcc cgt ggc atg gtc Ser Leu Trp Arg Phe Pro His Met Leu Glu Asn Ala Arg Gly Met Val 2020 2025 2030			6095

tcc caa ctc acc cag ttc ggt tcc acc ctc cag aac atc att gag agg Ser Gln Leu Thr Gln Phe Gly Ser Thr Leu Gln Asn Ile Ile Glu Arg 2035 2040 2045	6143
caa gat gct gag gct ctc aac gct ttg ctc cag aac cag gca gct gag Gln Asp Ala Glu Ala Leu Asn Ala Leu Leu Gln Asn Gln Ala Ala Glu 2050 2055 2060	6191
ttg atc ctc acc aac ttg tcc atc caa gac aag acc att gag gag ctt Leu Ile Leu Thr Asn Leu Ser Ile Gln Asp Lys Thr Ile Glu Glu Leu 2065 2070 2075	6239
gat gct gag aag aca gtc ctt gag aag agc aag gct ggt gcc caa tct Asp Ala Glu Lys Thr Val Leu Glu Lys Ser Lys Ala Gly Ala Gln Ser 2080 2085 2090 2095	6287
cgc ttc gac tcc tac ggc aag ctc tac gat gag aac atc aac gct ggt Arg Phe Asp Ser Tyr Gly Lys Leu Tyr Asp Glu Asn Ile Asn Ala Gly 2100 2105 2110	6335
gag aac cag gcc atg acc ctc agg gct tcc gca gct ggt ctc acc act Glu Asn Gln Ala Met Thr Leu Arg Ala Ser Ala Ala Gly Leu Thr Thr 2115 2120 2125	6383
gct gtc caa gcc tct cgc ttg gct ggt gca gct gct gac ctc gtt cca Ala Val Gln Ala Ser Arg Leu Ala Gly Ala Ala Ala Asp Leu Val Pro 2130 2135 2140	6431
aac atc ttc ggt ttc gct ggt ggt ggc tcc aga tgg ggt gcc att gct Asn Ile Phe Gly Phe Ala Gly Gly Ser Arg Trp Gly Ala Ile Ala 2145 2150 2155	6479
gag gct acc ggt tac gtc atg gag ttc tct gcc aac gtc atg aac act Glu Ala Thr Gly Tyr Val Met Glu Phe Ser Ala Asn Val Met Asn Thr 2160 2165 2170 2175	6527
gag gct gac aag atc agc caa tct gag acc tac aga agg cgc cgt caa Glu Ala Asp Lys Ile Ser Gln Ser Glu Thr Tyr Arg Arg Arg Gln 2180 2185 2190	6575
gag tgg gag atc caa agg aac aac gct gag gca gag ttg aag caa atc Glu Trp Glu Ile Gln Arg Asn Asn Ala Glu Ala Glu Leu Lys Gln Ile 2195 2200 2205	6623
gat gct caa ctc aag tcc ttg gct gtc aga agg gag gct gct gtc ctc Asp Ala Gln Leu Lys Ser Leu Ala Val Arg Arg Glu Ala Ala Val Leu 2210 2215 2220	6671
cag aag acc tcc ctc aag acc caa cag gag caa acc cag tcc cag ttg Gln Lys Thr Ser Leu Lys Thr Gln Gln Glu Gln Thr Gln Ser Gln Leu 2225 2230 2235	6719
gct ttc ctc caa agg aag ttc tcc aac cag gct ctc tac aac tgg ctc Ala Phe Leu Gln Arg Lys Phe Ser Asn Gln Ala Leu Tyr Asn Trp Leu 2240 2245 2250 2255	6767
aga ggc cgc ttg gct gcc atc tac ttc caa ttc tac gac ctt gct gtg Arg Gly Arg Leu Ala Ala Ile Tyr Phe Gln Phe Tyr Asp Leu Ala Val 2260 2265 2270	6815

gcc agg tgc ctc atg gct gag caa gcc tac cgc tgg gag ttg aac gat Ala Arg Cys Leu Met Ala Glu Gln Ala Tyr Arg Trp Glu Leu Asn Asp 2275 2280 2285	6863
gac tcc gcc agg ttc atc aag cca ggt gct tgg caa ggc acc tac gct Asp Ser Ala Arg Phe Ile Lys Pro Gly Ala Trp Gln Gly Thr Tyr Ala 2290 2295 2300	6911
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Lys Leu Gln Leu Thr Cys Pro Ala Glu Ile Ala Leu Tyr Pro Phe Asp		
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 Thr Phe Arg Glu Lys Thr Arg Gly Met Val Asn Trp Gly Glu Ala Lys
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 Arg Ile Tyr Glu Ile Ala Gln Ala Glu Gln Asp Arg Asn Leu Leu His
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gag aag agg atc ttc gcc tac gct aac cca ttg ctc aag aac gct gtc 239
Glu Lys Arg Ile Phe Ala Tyr Ala Asn Pro Leu Leu Lys Asn Ala Val
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Arg Leu Gly Thr Arg Gln Met Leu Gly Phe Ile Gln Gly Tyr Ser Asp
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Leu Phe Gly Asn Arg Ala Asp Asn Tyr Ala Ala Pro Gly Ser Val Ala
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  Ser Met Phe Ser Pro Ala Ala Tyr Leu Thr Glu Leu Tyr Arg Glu Ala
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Pro Asp Leu Ala Ser Leu Met Leu Ser Gln Lys Asn Met Asp Glu Glu
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 Ile Ser Thr Leu Ala Leu Ser Asn Glu Leu Cys Leu Ala Gly Ile Glu
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Arg Glu Ile Val His Glu Arg Asp Pro Gly Phe Arg His Leu Ser Gln			
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Gly Asp Ile Thr Thr Ala Gln Leu Met Ser Pro Ser Tyr Leu Ala Arg			
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Tyr Tyr Gly Val Ser Pro Glu Asp Ile Ala Tyr Val Thr Thr Ser Leu			
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Ser His Val Gly Tyr Ser Ser Asp Ile Leu Val Ile Pro Leu Val Asp			
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aac tac acc tcc cag acc aac tac att gag ttg tac cca caa ggt ggt	1055		
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Asp Asn Tyr Leu Ile Lys Tyr Asn Leu Ser Asn Ser Phe Gly Leu Asp			
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Ala His Asn Pro Tyr Pro Asp Met Val Ile Asn Gln Lys Tyr Glu Ser			
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caa gcc acc atc aag aga tct gac tct gac aac atc ctc tcc att ggt	1247		
Gln Ala Thr Ile Lys Arg Ser Asp Ser Asp Asn Ile Leu Ser Ile Gly			
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ctc caa agg tgg cac tct ggt tcc tac aac ttc gct gct gcc aac ttc	1295		

Leu Gln Arg Trp His Ser Gly Ser Tyr Asn Phe Ala Ala Ala Asn Phe			
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Lys Ile Asp Gln Tyr Ser Pro Lys Ala Phe Leu Leu Lys Met Asn Lys			
435	440	445	
gcc atc agg ctc ttg aag gcc act ggt ctc tcc ttc gcc acc ctt gag		1391	
Ala Ile Arg Leu Leu Lys Ala Thr Gly Leu Ser Phe Ala Thr Leu Glu			
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agg att gtg gac tct gtc aac tcc acc aag tcc atc act gtg gag gtc		1439	
Arg Ile Val Asp Ser Val Asn Ser Thr Lys Ser Ile Thr Val Glu Val			
465	470	475	
ctc aac aag gtc tac aga gtc aag ttc tac att gac cgc tac ggc atc		1487	
Leu Asn Lys Val Tyr Arg Val Lys Phe Tyr Ile Asp Arg Tyr Gly Ile			
480	485	490	495
tct gag gag act gct gcc atc ctt gcc aac atc aac atc tcc cag caa		1535	
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Pro Leu Asn Gly Ile Arg Tyr Glu Ile Ser Glu Asp Asn Ser Lys His			
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Leu Pro Asn Pro Asp Leu Asn Leu Lys Pro Asp Ser Thr Gly Asp Asp			
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Gln Arg Lys Ala Val Leu Lys Arg Ala Phe Gln Val Asn Ala Ser Glu			
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Lys Asn Asn Leu Glu Asn Leu Ser Asp Leu Tyr Leu Val Ser Leu Leu			
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Ala Gln Ile His Asn Leu Thr Ile Ala Glu Leu Asn Ile Leu Leu Val			
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Ile Cys Gly Tyr Gly Asp Thr Asn Ile Tyr Gln Ile Thr Asp Asp Asn			
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Leu Ala Lys Ile Val Glu Thr Leu Leu Trp Ile Thr Gln Trp Leu Lys			
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Val Tyr Tyr Pro Glu Asn Tyr Val Asp Pro Thr Gln Arg Ile Gly Gln	
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1650	1655	1660	
ttc aag gag aac gac tcc ttc gtc atc tac caa ggt gag ttg tct gag Phe Lys Glu Asn Asp Ser Phe Val Ile Tyr Gln Gly Glu Leu Ser Glu			5039
1665	1670	1675	
acc tcc caa act gtg gtc aag gtc ttc ctc tcc tac ttc att gag gcc Thr Ser Gln Thr Val Val Lys Val Phe Leu Ser Tyr Phe Ile Glu Ala			5087
1680	1685	1690	1695
acc ggt aac aag aac cac ctc tgg gtc agg gcc aag tac cag aag gag Thr Gly Asn Lys Asn His Leu Trp Val Arg Ala Lys Tyr Gln Lys Glu			5135
1700	1705	1710	
acc act gac aag atc ctc ttc gac agg act gat gag aag gac cca cat Thr Thr Asp Lys Ile Leu Phe Asp Arg Thr Asp Glu Lys Asp Pro His			5183
1715	1720	1725	
ggg tgg ttc ctc tct gat gac cac aag acc ttc tct ggt ctc agc tct Gly Trp Phe Leu Ser Asp Asp His Lys Thr Phe Ser Gly Leu Ser Ser			5231
1730	1735	1740	
gct caa gct ctc aag aac gac tct gag cca atg gac ttc tct ggt gcc Ala Gln Ala Leu Lys Asn Asp Ser Glu Pro Met Asp Phe Ser Gly Ala			5279
1745	1750	1755	
aac gct ctc tac ttc tgg gag ttg ttc tac tac act cca atg atg atg Asn Ala Leu Tyr Phe Trp Glu Leu Phe Tyr Tyr Thr Pro Met Met Met			5327
1760	1765	1770	1775
gct cac agg ctc ctt caa gag cag aac ttc gat gct gcc aac cac tgg Ala His Arg Leu Leu Gln Glu Gln Asn Phe Asp Ala Ala Asn His Trp			5375
1780	1785	1790	
ttc cgc tac gtc tgg agc cca tct ggt tac att gtg gat ggc aag att Phe Arg Tyr Val Trp Ser Pro Ser Gly Tyr Ile Val Asp Gly Lys Ile			5423
1795	1800	1805	
gcc atc tac cac tgg aac gtc agg cca ttg gag gag gac acc tcc tgg Ala Ile Tyr His Trp Asn Val Arg Pro Leu Glu Glu Asp Thr Ser Trp			5471
1810	1815	1820	
aac gct cag caa ctt gac tcc act gac cca gat gct gtg gct caa gat Asn Ala Gln Gln Leu Asp Ser Thr Asp Pro Asp Ala Val Ala Gln Asp			5519
1825	1830	1835	
gac cca atg cac tac aag gtg gcc acc ttc atg gcc acc ttg gac ctt Asp Pro Met His Tyr Lys Val Ala Thr Phe Met Ala Thr Leu Asp Leu			5567
1840	1845	1850	1855
ctc atg gcc aga ggt gat gct gcc tac cgc caa ttg gag agg gac acc Leu Met Ala Arg Gly Asp Ala Ala Tyr Arg Gln Leu Glu Arg Asp Thr			5615
1860	1865	1870	
ttg gct gag gcc aag atg tgg tac acc caa gct ctc aac ttg ctg ggt Leu Ala Glu Ala Lys Met Trp Tyr Thr Gln Ala Leu Asn Leu Leu Gly			5663
1875	1880	1885	

gat gag cca caa gtc atg ctc tcc aca acc tgg gcc aac cca acc ttg Asp Glu Pro Gln Val Met Leu Ser Thr Thr Trp Ala Asn Pro Thr Leu 1890 1895 1900	5711
ggc aac gct gcc tcc aag acc aca caa cag gtc agg caa cag gtc ctc Gly Asn Ala Ala Ser Lys Thr Thr Gln Gln Val Arg Gln Gln Val Leu 1905 1910 1915	5759
acc caa ctc agg ctc aac tct aga gtc aag act cca ctc ttg ggc act Thr Gln Leu Arg Leu Asn Ser Arg Val Lys Thr Pro Leu Leu Gly Thr 1920 1925 1930 1935	5807
gcc aac tcc ctc act gct ctc ctc cca caa gag aac tcc aaa ctt Ala Asn Ser Leu Thr Ala Leu Phe Leu Pro Gln Glu Asn Ser Lys Leu 1940 1945 1950	5855
aag ggt tac tgg agg acc ctt gct caa cgc atg ttc aac ctc agg cac Lys Gly Tyr Trp Arg Thr Leu Ala Gln Arg Met Phe Asn Leu Arg His 1955 1960 1965	5903
aac ctc tcc att gat ggt caa cca ctc tcc ttg cca ctc tac gct aag Asn Leu Ser Ile Asp Gly Gln Pro Leu Ser Leu Pro Leu Tyr Ala Lys 1970 1975 1980	5951
cca gct gac cca aag gct ctc ctt tcc gct gtc tcc gca tcc caa Pro Ala Asp Pro Lys Ala Leu Leu Ser Ala Ala Val Ser Ala Ser Gln 1985 1990 1995	5999
ggt ggt gct gac ctc cca aag gct cca ctc acc atc cac agg ttc cca Gly Gly Ala Asp Leu Pro Lys Ala Pro Leu Thr Ile His Arg Phe Pro 2000 2005 2010 2015	6047
caa atg ttg gag ggt gcc cgt ggt ctt gtc aac cag ctc atc caa ttc Gln Met Leu Glu Gly Ala Arg Gly Leu Val Asn Gln Leu Ile Gln Phe 2020 2025 2030	6095
ggt tcc tct ctc ctt ggt tac tct gag agg caa gat gct gag gcc atg Gly Ser Ser Leu Leu Gly Tyr Ser Glu Arg Gln Asp Ala Glu Ala Met 2035 2040 2045	6143
tcc caa ctc ttg caa acc cag gct tct gag ttg atc ctc acc tcc atc Ser Gln Leu Leu Gln Thr Gln Ala Ser Glu Leu Ile Leu Thr Ser Ile 2050 2055 2060	6191
agg atg caa gac aac cag ctt gct gag ttg gac tct gag aag act gct Arg Met Gln Asp Asn Gln Leu Ala Glu Leu Asp Ser Glu Lys Thr Ala 2065 2070 2075	6239
ctc caa gtc tcc ctt gct ggt gtc caa cag agg ttc gac agc tac tcc Leu Gln Val Ser Leu Ala Gly Val Gln Gln Arg Phe Asp Ser Tyr Ser 2080 2085 2090 2095	6287
caa ctc tac gag gag aac atc aac gct ggt gag caa agg gct ttg gct Gln Leu Tyr Glu Glu Asn Ile Asn Ala Gly Glu Gln Arg Ala Leu Ala 2100 2105 2110	6335
ctc agg tct gag tct gcc att gag tcc caa ggt gct caa atc tcc cgc Leu Arg Ser Glu Ser Ala Ile Glu Ser Gln Gly Ala Gln Ile Ser Arg 2115 2120 2125	6383

atg gct ggt gct ggc gtg gac atg gct cca aac atc ttc ggt ctt gct 6431
 Met Ala Gly Ala Gly Val Asp Met Ala Pro Asn Ile Phe Gly Leu Ala
 2130 2135 2140

gat ggt ggc atg cac tac ggt gcc att gct tac gcc att gct gat ggc 6479
 Asp Gly Gly Met His Tyr Gly Ala Ile Ala Tyr Ala Ile Ala Asp Gly
 2145 2150 2155

att gag ctt tct gct tct gcc aag atg gtt gat gct gag aag gtg gct 6527
 Ile Glu Leu Ser Ala Ser Ala Lys Met Val Asp Ala Glu Lys Val Ala
 2160 2165 2170 2175

caa tct gaa atc tac cgt cgc aga cgc caa gaa tgg aag atc caa agg 6575
 Gln Ser Glu Ile Tyr Arg Arg Arg Gln Glu Trp Lys Ile Gln Arg
 2180 2185 2190

gac aac gct caa gct gag atc aac cag ctc aac gct caa ctt gag tcc 6623
 Asp Asn Ala Gln Ala Glu Ile Asn Gln Leu Asn Ala Gln Leu Glu Ser
 2195 2200 2205

ctc agc atc agg cgt gag gct gct gag atg cag aag gag tac ctc aag 6671
 Leu Ser Ile Arg Arg Glu Ala Ala Glu Met Gln Lys Glu Tyr Leu Lys
 2210 2215 2220

acc caa cag gct caa gct cag gct caa ctc acc ttc ctc agg tcc aag 6719
 Thr Gln Gln Ala Gln Ala Gln Leu Thr Phe Leu Arg Ser Lys
 2225 2230 2235

ttc tcc aac cag gct ctc tac tcc tgg ctc aga ggc cgc ctc tct ggc 6767
 Phe Ser Asn Gln Ala Leu Tyr Ser Trp Leu Arg Gly Arg Leu Ser Gly
 2240 2245 2250 2255

atc tac ttc caa ttc tac gac ttg gct gtc tcc cgc tgc ctc atg gct 6815
 Ile Tyr Phe Gln Phe Tyr Asp Leu Ala Val Ser Arg Cys Leu Met Ala
 2260 2265 2270

gag caa tcc tac caa tgg gag gcc aac gac aac agc atc tcc ttc gtc 6863
 Glu Gln Ser Tyr Gln Trp Glu Ala Asn Asp Asn Ser Ile Ser Phe Val
 2275 2280 2285

aag cca ggt gct tgg caa ggc acc tac gct ggt ctc ctt tgc ggt gag 6911
 Lys Pro Gly Ala Trp Gln Gly Thr Tyr Ala Gly Leu Leu Cys Gly Glu
 2290 2295 2300

gct ctc atc cag aac ttg gct caa atg gag gag gct tac ctc aag tgg 6959
 Ala Leu Ile Gln Asn Leu Ala Gln Met Glu Glu Ala Tyr Leu Lys Trp
 2305 2310 2315

gag tcc aga gct ttg gag gta gag agg act gtc tcc ctt gct gta gtc 7007
 Glu Ser Arg Ala Leu Glu Val Glu Arg Thr Val Ser Leu Ala Val Val
 2320 2325 2330 2335

tac gac tcc ttg gag ggc aac gac agg ttc aac ctt gct gag caa atc 7055
 Tyr Asp Ser Leu Glu Gly Asn Asp Arg Phe Asn Leu Ala Glu Gln Ile
 2340 2345 2350

cca gct ctc ttg gac aag ggt gag ggc act gct ggc acc aag gag aac 7103
 Pro Ala Leu Leu Asp Lys Gly Glu Gly Thr Ala Gly Thr Lys Glu Asn
 2355 2360 2365

ggt ctc tcc ttg gcc aac gcc atc ctc tct gct tct gtc aag ctc tct 7151

Gly Leu Ser Leu Ala Asn Ala Ile Leu Ser Ala Ser Val Lys Leu Ser				
2370	2375	2380		
gac ctc aag ttg ggt act gac tac cca gac tcc att gtg ggt tcc aac				7199
Asp Leu Lys Leu Gly Thr Asp Tyr Pro Asp Ser Ile Val Gly Ser Asn				
2385	2390	2395		
aag gtc aga agg atc aag caa atc tct gtc tcc ctc cca gct ttg gtg				7247
Lys Val Arg Arg Ile Lys Gln Ile Ser Val Ser Leu Pro Ala Leu Val				
2400	2405	2410	2415	
ggt cca tac caa gat gtc caa gcc atg ctc tcc tac ggt ggc tcc acc				7295
Gly Pro Tyr Gln Asp Val Gln Ala Met Leu Ser Tyr Gly Gly Ser Thr				
2420	2425	2430		
caa ctc cca aag ggt tgc tct gct ttg gct gtc tcc cac ggc acc aac				7343
Gln Leu Pro Lys Gly Cys Ser Ala Leu Ala Val Ser His Gly Thr Asn				
2435	2440	2445		
gac tct ggt caa ttc caa ctt gac ttc aac gat ggc aag tac ctc cca				7391
Asp Ser Gly Gln Phe Gln Leu Asp Phe Asn Asp Gly Lys Tyr Leu Pro				
2450	2455	2460		
ttc gaa ggc att gct ttg gat gac caa ggc acc ctc aac ctc caa ttc				7439
Phe Glu Gly Ile Ala Leu Asp Asp Gln Gly Thr Leu Asn Leu Gln Phe				
2465	2470	2475		
cca aac gcc act gac aag cag aag gcc atc ctc caa acc atg tct gac				7487
Pro Asn Ala Thr Asp Lys Gln Lys Ala Ile Leu Gln Thr Met Ser Asp				
2480	2485	2490	2495	
atc atc ctc cac atc agg tac acc atc agg tgagctcgag aggcctgcgg				7537
Ile Ile Leu His Ile Arg Tyr Thr Ile Arg				
2500	2505			
ccgc				7541
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encoding ER signal from 15 kDa zein from Black				
Mexican Sweet maize				
<220>				
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Met Ala Lys Met Val Ile Val Leu Val Val Cys Leu Ala Leu Ser Ala				
1	5	10	15	
gcc tgt gct tca gcc				63
Ala Cys Ala Ser Ala				
20				

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 reticulum signal peptide

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gct gcc tgt gct tca gcc atg aac gag tcc gtc aag gag atc cca gac	96
Ala Ala Cys Ala Ser Ala Met Asn Glu Ser Val Lys Glu Ile Pro Asp	
20 25 30	
gtc ctc aag tcc caa tgc ggt ttc aac tgc ctc act gac atc tcc cac	144
Val Leu Lys Ser Gln Cys Gly Phe Asn Cys Leu Thr Asp Ile Ser His	
35 40 45	
agc tcc ttc aac gag ttc aga caa caa gtc tct gag cac ctc tcc tgg	192
Ser Ser Phe Asn Glu Phe Arg Gln Gln Val Ser Glu His Leu Ser Trp	
50 55 60	
tcc gag acc cat gac ctc tac cat gac gct cag caa gct cag aag gac	240
Ser Glu Thr His Asp Leu Tyr His Asp Ala Gln Gln Ala Gln Lys Asp	
65 70 75	
aac agg ctc tac gag gct agg atc ctc aag agg gct aac cca caa ctc	288
Asn Arg Leu Tyr Glu Ala Arg Ile Leu Lys Arg Ala Asn Pro Gln Leu	
80 85 90 95	
cag aac gct gtc cac ctc gcc atc ttg gct cca aac gct gag ttg att	336
Gln Asn Ala Val His Leu Ala Ile Leu Ala Pro Asn Ala Glu Leu Ile	
100 105 110	
ggc tac aac aac cag ttc tct ggc aga gct agc cag tac gtg gct cct	384
Gly Tyr Asn Asn Gln Phe Ser Gly Arg Ala Ser Gln Tyr Val Ala Pro	
115 120 125	
ggc aca gtc tcc tcc atg ttc agc cca gcc gct tac ctc act gag ttg	432
Gly Thr Val Ser Ser Met Phe Ser Pro Ala Ala Tyr Leu Thr Glu Leu	
130 135 140	
tac cgc gag gct agg aac ctt cat gct tct gac tcc gtc tac tac ttg	480
Tyr Arg Glu Ala Arg Asn Leu His Ala Ser Asp Ser Val Tyr Tyr Leu	
145 150 155	
gac aca cgc aga cca gac ctc aag agc atg gcc ctc agc caa cag aac	528
Asp Thr Arg Arg Pro Asp Leu Lys Ser Met Ala Leu Ser Gln Gln Asn	
160 165 170 175	
atg gac att gag ttg tcc acc ctc tcc ttg agc aac gag ctt ctc ttg	576

Met Asp Ile Glu Leu Ser Thr Leu Ser Leu Ser Asn Glu Leu Leu Leu			
180	185	190	
gag tcc atc aag act gag agc aag ttg gag aac tac acc aag gtc atg 624			
Glu Ser Ile Lys Thr Glu Ser Lys Leu Glu Asn Tyr Thr Lys Val Met			
195	200	205	
gag atg ctc tcc acc ttc aga cca agc ggt gca act cca tac cat gat 672			
Glu Met Leu Ser Thr Phe Arg Pro Ser Gly Ala Thr Pro Tyr His Asp			
210	215	220	
gcc tac gag aac gtc agg gag gtc atc caa ctt caa gac cct ggt ctt 720			
Ala Tyr Glu Asn Val Arg Glu Val Ile Gln Leu Gln Asp Pro Gly Leu			
225	230	235	
gag caa ctc aac gct tct cca gcc att gct ggt ttg atg cac caa gca 768			
Glu Gln Leu Asn Ala Ser Pro Ala Ile Ala Gly Leu Met His Gln Ala			
240	245	250	255
tcc ttg ctc ggt atc aac gcc tcc atc tct cct gag ttg ttc aac atc 816			
Ser Leu Leu Gly Ile Asn Ala Ser Ile Ser Pro Glu Leu Phe Asn Ile			
260	265	270	
ttg act gag gag atc act gag ggc aac gct gag gag ttg tac aag aag 864			
Leu Thr Glu Glu Ile Thr Glu Gly Asn Ala Glu Glu Leu Tyr Lys Lys			
275	280	285	
aac ttc ggc aac att gag cca gcc tct ctt gca atg cct gag tac ctc 912			
Asn Phe Gly Asn Ile Glu Pro Ala Ser Leu Ala Met Pro Glu Tyr Leu			
290	295	300	
aag agg tac tac aac ttg tct gat gag gag ctt tct caa ttc att ggc 960			
Lys Arg Tyr Tyr Asn Leu Ser Asp Glu Glu Leu Ser Gln Phe Ile Gly			
305	310	315	
aag gct tcc aac ttc ggt caa cag gag tac agc aac aac cag ctc atc 1008			
Lys Ala Ser Asn Phe Gly Gln Gln Glu Tyr Ser Asn Asn Gln Leu Ile			
320	325	330	335
act cca gtt gtg aac tcc tct gat ggc act gtg aag gtc tac cgc atc 1056			
Thr Pro Val Val Asn Ser Ser Asp Gly Thr Val Lys Val Tyr Arg Ile			
340	345	350	
aca cgt gag tac acc aca aac gcc tac caa atg gat gtt gag ttg ttc 1104			
Thr Arg Glu Tyr Thr Asn Ala Tyr Gln Met Asp Val Glu Leu Phe			
355	360	365	
cca ttc ggt ggt gag aac tac aga ctt gac tac aag ttc aag aac ttc 1152			
Pro Phe Gly Gly Glu Asn Tyr Arg Leu Asp Tyr Lys Phe Lys Asn Phe			
370	375	380	
tac aac gcc tcc tac ctc tcc atc aag ttg aac gac aag agg gag ctt 1200			
Tyr Asn Ala Ser Tyr Leu Ser Ile Lys Leu Asn Asp Lys Arg Glu Leu			
385	390	395	
gtc agg act gag ggt gct cca gtg aac att gag tac tct gcc aac 1248			
Val Arg Thr Glu Gly Ala Pro Gln Val Asn Ile Glu Tyr Ser Ala Asn			
400	405	410	415
atc acc ctc aac aca gct gac atc tct caa cca ttc gag att ggt ttg 1296			
Ile Thr Leu Asn Thr Ala Asp Ile Ser Gln Pro Phe Glu Ile Gly Leu			

420	425	430	
acc aga gtc ctt ccc tct ggc tcc tgg gcc tac gct gca gcc aag ttc Thr Arg Val Leu Pro Ser Gly Ser Trp Ala Tyr Ala Ala Ala Lys Phe			1344
435	440	445	
act gtt gag gag tac aac cag tac tct ttc ctc ttg aag ctc aac aag Thr Val Glu Glu Tyr Asn Gln Tyr Ser Phe Leu Leu Lys Leu Asn Lys			1392
450	455	460	
gca att cgt ctc agc aga gcc act gag ttg tct ccc acc atc ttg gag Ala Ile Arg Leu Ser Arg Ala Thr Glu Leu Ser Pro Thr Ile Leu Glu			1440
465	470	475	
ggc att gtg agg tct gtc aac ctt caa ctt gac atc aac act gat gtg Gly Ile Val Arg Ser Val Asn Leu Gln Leu Asp Ile Asn Thr Asp Val			1488
480	485	490	495
ctt ggc aag gtc ttc ctc acc aag tac tac atg caa cgc tac gcc atc Leu Gly Lys Val Phe Leu Thr Lys Tyr Tyr Met Gln Arg Tyr Ala Ile			1536
500	505	510	
cat gct gag act gca ctc atc ctc tgc aac gca ccc atc tct caa cgc His Ala Glu Thr Ala Leu Ile Leu Cys Asn Ala Pro Ile Ser Gln Arg			1584
515	520	525	
tcc tac gac aac cag cct tcc cag ttc gac agg ctc ttc aac act cct Ser Tyr Asp Asn Gln Pro Ser Gln Phe Asp Arg Leu Phe Asn Thr Pro			1632
530	535	540	
ctc ttg aac ggc cag tac ttc tcc act ggt gat gag gag att gac ctc Leu Leu Asn Gly Gln Tyr Phe Ser Thr Gly Asp Glu Glu Ile Asp Leu			1680
545	550	555	
aac tct ggc tcc aca ggt gac tgg aga aag acc atc ttg aag agg gcc Asn Ser Gly Ser Thr Gly Asp Trp Arg Lys Thr Ile Leu Lys Arg Ala			1728
560	565	570	575
ttc aac att gat gat gtc tct ctc cgt ctc ttg aag atc aca gat Phe Asn Ile Asp Asp Val Ser Leu Phe Arg Leu Leu Lys Ile Thr Asp			1776
580	585	590	
cac gac aac aag gat ggc aag atc aag aac aac ttg aag aac ctt tcc His Asp Asn Lys Asp Gly Lys Ile Lys Asn Asn Leu Lys Asn Leu Ser			1824
595	600	605	
aac ctc tac att ggc aag ttg ctt gca gac atc cac caa ctc acc att Asn Leu Tyr Ile Gly Lys Leu Leu Ala Asp Ile His Gln Leu Thr Ile			1872
610	615	620	
gat gag ttg gac ctc ttg ctc att gca gtc ggt gag ggc aag acc aac Asp Glu Leu Asp Leu Leu Ile Ala Val Gly Glu Gly Lys Thr Asn			1920
625	630	635	
ctc tct gca atc tct gac aag cag ttg gca acc ctc atc agg aag ttg Leu Ser Ala Ile Ser Asp Lys Gln Leu Ala Thr Leu Ile Arg Lys Leu			1968
640	645	650	655
aac acc atc acc tcc tgg ctt cac acc cag aag tgg tct gtc ttc caa Asn Thr Ile Thr Ser Trp Leu His Thr Gln Lys Trp Ser Val Phe Gln			2016
660	665	670	

ctc ttc atc atg acc agc acc tcc tac aac aag acc ctc act cct gag Leu Phe Ile Met Thr Ser Thr Ser Tyr Asn Lys Thr Leu Thr Pro Glu 675 680 685	2064
atc aag aac ctc ttg gac aca gtc tac cac ggt ctc caa ggc ttc gac Ile Lys Asn Leu Leu Asp Thr Val Tyr His Gly Leu Gln Gly Phe Asp 690 695 700	2112
aag gac aag gct gac ttg ctt cat gtc atg gct ccc tac att gca gcc Lys Asp Lys Ala Asp Leu Leu His Val Met Ala Pro Tyr Ile Ala Ala 705 710 715	2160
acc ctc caa ctc tcc tct gag aac gtg gct cac tct gtc ttg ctc tgg Thr Leu Gln Leu Ser Ser Glu Asn Val Ala His Ser Val Leu Leu Trp 720 725 730 735	2208
gct gac aag ctc caa cct ggt gat ggt gcc atg act gct gag aag ttc Ala Asp Lys Leu Gln Pro Gly Asp Gly Ala Met Thr Ala Glu Lys Phe 740 745 750	2256
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gag act caa gag cac att gtg caa tac tgc cag gct ctt gca cag ttg Glu Thr Gln Glu His Ile Val Gln Tyr Cys Gln Ala Leu Ala Gln Leu 770 775 780	2352
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gct cat gat gct ctc tcc ctc atc atg ttg acc agg ttc gct gac tgg Ala His Asp Ala Leu Ser Leu Ile Met Leu Thr Arg Phe Ala Asp Trp 820 825 830	2496
gtc aac gct ctt ggt gag aag gct tcc tct gtc ttg gct gcc ttc gag Val Asn Ala Leu Gly Glu Lys Ala Ser Ser Val Leu Ala Ala Phe Glu 835 840 845	2544
gcc aac tcc ctc act gct gag caa ctt gct gat gcc atg aac ctt gat Ala Asn Ser Leu Thr Ala Glu Gln Leu Ala Asp Ala Met Asn Leu Asp 850 855 860	2592
gcc aac ctc ttg ctc caa gct tcc att caa gct cag aac cac caa cac Ala Asn Leu Leu Gln Ala Ser Ile Gln Ala Gln Asn His Gln His 865 870 875	2640
ctc cca cct gtc act cca gag aac gct ttc tcc tgc tgg acc tcc atc Leu Pro Pro Val Thr Pro Glu Asn Ala Phe Ser Cys Trp Thr Ser Ile 880 885 890 895	2688
aac acc atc ctc caa tgg gtc aac gtg gct cag caa ctc aac gtg gct Asn Thr Ile Leu Gln Trp Val Asn Val Ala Gln Gln Leu Asn Val Ala 900 905 910	2736

cca caa ggt gtc tct gct ttg gtc ggt ctt gac tac atc cag tcc atg	2784
Pro Gln Gly Val Ser Ala Leu Val Gly Leu Asp Tyr Ile Gln Ser Met	
915 920 925	
aag gag aca cca acc tac gct caa tgg gag aac gca gct ggt gtc ttg	2832
Lys Glu Thr Pro Thr Tyr Ala Gln Trp Glu Asn Ala Ala Gly Val Leu	
930 935 940	
act gct ggt ctc aac tcc caa cag gcc aac acc ctc cat gct ttc ttg	2880
Thr Ala Gly Leu Asn Ser Gln Gln Ala Asn Thr Leu His Ala Phe Leu	
945 950 955	
gat gag tct cgc tct gct gcc ctc acc tac tac atc agg caa gtc	2928
Asp Glu Ser Arg Ser Ala Ala Leu Ser Thr Tyr Tyr Ile Arg Gln Val	
960 965 970 975	
gcc aag gca gct gct gcc atc aag tct cgc gat gac ctc tac caa tac	2976
Ala Lys Ala Ala Ala Ile Lys Ser Arg Asp Asp Leu Tyr Gln Tyr	
980 985 990	
ctc ctc att gac aac cag gtc tct gct gcc atc aag acc acc agg atc	3024
Leu Leu Ile Asp Asn Gln Val Ser Ala Ala Ile Lys Thr Thr Arg Ile	
995 1000 1005	
gct gag gcc atc gct tcc atc caa ctc tac gtc aac cgc gct ctt gag	3072
Ala Glu Ala Ile Ala Ser Ile Gln Leu Tyr Val Asn Arg Ala Leu Glu	
1010 1015 1020	
aac gtt gag gag aac gcc aac tct ggt gtc atc tct cgc caa ttc ttc	3120
Asn Val Glu Glu Asn Ala Asn Ser Gly Val Ile Ser Arg Gln Phe Phe	
1025 1030 1035	
atc gac tgg gac aag tac aac aag agg tac tcc acc tgg gct ggt gtc	3168
Ile Asp Trp Asp Lys Tyr Asn Lys Arg Tyr Ser Thr Trp Ala Gly Val	
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tct caa ctt gtc tac tac cca gag aac tac att gac cca acc atg agg	3216
Ser Gln Leu Val Tyr Tyr Pro Glu Asn Tyr Ile Asp Pro Thr Met Arg	
1060 1065 1070	
att ggt cag acc aag atg atg gat gct ctc ttg caa tct gtc tcc caa	3264
Ile Gly Gln Thr Lys Met Met Asp Ala Leu Leu Gln Ser Val Ser Gln	
1075 1080 1085	
agc caa ctc aac gct gac act gtg gag gat gcc ttc atg agc tac ctc	3312
Ser Gln Leu Asn Ala Asp Thr Val Glu Asp Ala Phe Met Ser Tyr Leu	
1090 1095 1100	
acc tcc ttc gag caa gtt gcc aac ctc aag gtc atc tct gct tac cat	3360
Thr Ser Phe Glu Gln Val Ala Asn Leu Lys Val Ile Ser Ala Tyr His	
1105 1110 1115	
gac aac atc aac aac gac caa ggt ctc acc tac ttc att ggt ctc tct	3408
Asp Asn Ile Asn Asn Asp Gln Gly Leu Thr Tyr Phe Ile Gly Leu Ser	
1120 1125 1130 1135	
gag act gat gct ggt gag tac tac tgg aga tcc gtg gac cac agc aag	3456
Glu Thr Asp Ala Gly Glu Tyr Tyr Trp Arg Ser Val Asp His Ser Lys	
1140 1145 1150	
ttc aac gat ggc aag ttc gct gca aac gct tgg tct gag tgg cac aag	3504

Phe Asn Asp Gly Lys Phe Ala Ala Asn Ala Trp Ser Glu Trp His Lys			
1155	1160	1165	
att gac tgc cct atc aac cca tac aag tcc acc atc aga cct gtc atc			3552
Ile Asp Cys Pro Ile Asn Pro Tyr Lys Ser Thr Ile Arg Pro Val Ile			
1170	1175	1180	
tac aag agc cgc ctc tac ttg ctc tgg ctt gag cag aag gag atc acc			3600
Tyr Lys Ser Arg Leu Tyr Leu Leu Trp Leu Glu Gln Lys Glu Ile Thr			
1185	1190	1195	
aag caa act ggc aac tcc aag gat ggt tac caa act gag act gac tac			3648
Lys Gln Thr Gly Asn Ser Lys Asp Gly Tyr Gln Thr Glu Thr Asp Tyr			
1200	1205	1210	1215
cgc tac gag ttg aag ttg gct cac atc cgc tac gat ggt acc tgg aac			3696
Arg Tyr Glu Leu Lys Leu Ala His Ile Arg Tyr Asp Gly Thr Trp Asn			
1220	1225	1230	
act cca atc acc ttc gat gtc aac aag aag atc agc gag ttg aag ttg			3744
Thr Pro Ile Thr Phe Asp Val Asn Lys Lys Ile Ser Glu Leu Lys Leu			
1235	1240	1245	
gag aag aac cgt gct cct ggt ctc tac tgc gct ggt tac caa ggt gag			3792
Glu Lys Asn Arg Ala Pro Gly Leu Tyr Cys Ala Gly Tyr Gln Gly Glu			
1250	1255	1260	
gac acc ctc ttg gtc atg ttc tac aac cag caa gac acc ctt gac tcc			3840
Asp Thr Leu Leu Val Met Phe Tyr Asn Gln Gln Asp Thr Leu Asp Ser			
1265	1270	1275	
tac aag aac gct tcc atg caa ggt ctc tac atc ttc gct gac atg gct			3888
Tyr Lys Asn Ala Ser Met Gln Gly Leu Tyr Ile Phe Ala Asp Met Ala			
1280	1285	1290	1295
tcc aag gac atg act cca gag caa agc aac gtc tac cgt gac aac tcc			3936
Ser Lys Asp Met Thr Pro Glu Gln Ser Asn Val Tyr Arg Asp Ash Ser			
1300	1305	1310	
tac caa cag ttc gac acc aac gtc agg cgt gtc aac aac aga tac			3984
Tyr Gln Gln Phe Asp Thr Asn Asn Val Arg Arg Val Asn Asn Arg Tyr			
1315	1320	1325	
gct gag gac tac gag atc cca agc tct gtc agc tct cgc aag gac tac			4032
Ala Glu Asp Tyr Glu Ile Pro Ser Ser Val Ser Ser Arg Lys Asp Tyr			
1330	1335	1340	
ggc tgg ggt gac tac tac ctc agc atg gtg tac aac ggt gac atc cca			4080
Gly Trp Gly Asp Tyr Tyr Leu Ser Met Val Tyr Asn Gly Asp Ile Pro			
1345	1350	1355	
acc atc aac tac aag gct gcc tct tcc gac ctc aaa atc tac atc agc			4128
Thr Ile Asn Tyr Lys Ala Ala Ser Ser Asp Leu Lys Ile Tyr Ile Ser			
1360	1365	1370	1375
cca aag ctc agg atc atc cac aac ggc tac gag ggt cag aag agg aac			4176
Pro Lys Leu Arg Ile Ile His Asn Gly Tyr Glu Gly Gln Lys Arg Asn			
1380	1385	1390	
cag tgc aac ttg atg aac aag tac ggc aag ttg ggt gac aag ttc att			4224
Gln Cys Asn Leu Met Asn Lys Tyr Gly Lys Leu Gly Asp Lys Phe Ile			

1395	1400	1405	
gtc tac acc tct ctt ggt gtc aac cca aac aac agc tcc aac aag ctc Val Tyr Thr Ser Leu Gly Val Asn Pro Asn Asn Ser Ser Asn Lys Leu			4272
1410	1415	1420	
atg ttc tac cca gtc tac caa tac tct ggc aac acc tct ggt ctc aac Met Phe Tyr Pro Val Tyr Gln Tyr Ser Gly Asn Thr Ser Gly Leu Asn			4320
1425	1430	1435	
cag ggt aga ctc ttg ttc cac agg gac acc acc tac cca agc aag gtg Gln Gly Arg Leu Leu Phe His Arg Asp Thr Thr Tyr Pro Ser Lys Val			4368
1440	1445	1450	1455
gag gct tgg att cct ggt gcc aag agg tcc ctc acc aac cag aac gct Glu Ala Trp Ile Pro Gly Ala Lys Arg Ser Leu Thr Asn Gln Asn Ala			4416
1460	1465	1470	
gcc att ggt gat gac tac gcc aca gac tcc ctc aac aag cct gat gac Ala Ile Gly Asp Asp Tyr Ala Thr Asp Ser Leu Asn Lys Pro Asp Asp			4464
1475	1480	1485	
ctc aag cag tac atc ttc atg act gac tcc aag ggc aca gcc act gat Leu Lys Gln Tyr Ile Phe Met Thr Asp Ser Lys Gly Thr Ala Thr Asp			4512
1490	1495	1500	
gtc tct ggt cca gtg gag atc aac act gca atc agc cca gcc aag gtc Val Ser Gly Pro Val Glu Ile Asn Thr Ala Ile Ser Pro Ala Lys Val			4560
1505	1510	1515	
caa atc att gtc aag gct ggt ggc aag gag caa acc ttc aca gct gac Gln Ile Ile Val Lys Ala Gly Gly Lys Glu Gln Thr Phe Thr Ala Asp			4608
1520	1525	1530	1535
aag gat gtc tcc atc cag cca agc cca tcc ttc gat gag atg aac tac Lys Asp Val Ser Ile Gln Pro Ser Pro Ser Phe Asp Glu Met Asn Tyr			4656
1540	1545	1550	
caa ttc aac gct ctt gag att gat ggt tct ggc ctc aac ttc atc aac Gln Phe Asn Ala Leu Glu Ile Asp Gly Ser Gly Leu Asn Phe Ile Asn			4704
1555	1560	1565	
aac tct gct tcc att gat gtc acc ttc act gcc ttc gct gag gat ggc Asn Ser Ala Ser Ile Asp Val Thr Phe Thr Ala Phe Ala Glu Asp Gly			4752
1570	1575	1580	
cgc aag ttg ggt tac gag agc ttc tcc atc cca gtc acc ctt aag gtt Arg Lys Leu Gly Tyr Glu Ser Phe Ser Ile Pro Val Thr Leu Lys Val			4800
1585	1590	1595	
tcc act gac aac gca ctc acc ctt cat cac aac gag aac ggt gct cag Ser Thr Asp Asn Ala Leu Thr Leu His His Asn Glu Asn Gly Ala Gln			4848
1600	1605	1610	1615
tac atg caa tgg caa agc tac cgc acc agg ttg aac acc ctc ttc gca Tyr Met Gln Trp Gln Ser Tyr Arg Thr Arg Leu Asn Thr Leu Phe Ala			4896
1620	1625	1630	
agg caa ctt gtg gcc cgt gcc acc aca ggc att gac acc atc ctc agc Arg Gln Leu Val Ala Arg Ala Thr Thr Gly Ile Asp Thr Ile Leu Ser			4944
1635	1640	1645	

atg gag acc cag aac atc caa gag cca cag ttg ggc aag ggt ttc tac Met Glu Thr Gln Asn Ile Gln Glu Pro Gln Leu Gly Lys Gly Phe Tyr 1650 1655 1660	4992
gcc acc ttc gtc atc cca cct tac aac ctc agc act cat ggt gat gag Ala Thr Phe Val Ile Pro Pro Tyr Asn Leu Ser Thr His Gly Asp Glu 1665 1670 1675	5040
agg tgg ttc aag ctc tac atc aag cac gtg gtt gac aac aac tcc cac Arg Trp Phe Lys Leu Tyr Ile Lys His Val Val Asp Asn Asn Ser His 1680 1685 1690 1695	5088
atc atc tac tct ggt caa ctc act gac acc aac atc aac atc acc ctc Ile Ile Tyr Ser Gly Gln Leu Thr Asp Thr Asn Ile Asn Ile Thr Leu 1700 1705 1710	5136
ttc atc cca ctt gac gat gtc cca ctc aac cag gac tac cat gcc aag Phe Ile Pro Leu Asp Asp Val Pro Leu Asn Gln Asp Tyr His Ala Lys 1715 1720 1725	5184
gtc tac atg acc ttc aag aag tct cca tct gat ggc acc tgg tgg ggt Val Tyr Met Thr Phe Lys Lys Ser Pro Ser Asp Gly Thr Trp Trp Gly 1730 1735 1740	5232
cca cac ttc gtc cgt gat gac aag ggc atc gtc acc atc aac cca aag Pro His Phe Val Arg Asp Asp Lys Gly Ile Val Thr Ile Asn Pro Lys 1745 1750 1755	5280
tcc atc ctc acc cac ttc gag tct gtc aac gtt ctc aac aac atc tcc Ser Ile Leu Thr His Phe Glu Ser Val Asn Val Leu Asn Asn Ile Ser 1760 1765 1770 1775	5328
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ttg ttc tac tac aca cca atg ctt gtg gct caa agg ttg ctc cat gag Leu Phe Tyr Tyr Pro Met Leu Val Ala Gln Arg Leu Leu His Glu 1795 1800 1805	5424
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tct ggt tac att gtg cat ggt caa atc cag aac tac caa tgg aac gtc Ser Gly Tyr Ile Val His Gly Gln Ile Gln Asn Tyr Gln Trp Asn Val 1825 1830 1835	5520
agg cca ttg ctt gag gac acc tcc tgg aac tct gac cca ctt gac tct Arg Pro Leu Leu Glu Asp Thr Ser Trp Asn Ser Asp Pro Leu Asp Ser 1840 1845 1850 1855	5568
gtg gac cct gat gct gtg gct caa cat gac cca atg cac tac aag gtc Val Asp Pro Asp Ala Val Ala Gln His Asp Pro Met His Tyr Lys Val 1860 1865 1870	5616
tcc acc ttc atg agg acc ttg gac ctc ttg att gcc aga ggt gac cat Ser Thr Phe Met Arg Thr Leu Asp Leu Leu Ile Ala Arg Gly Asp His 1875 1880 1885	5664

gct tac cgc caa ttg gag agg gac acc ctc aac gag gca aag atg tgg Ala Tyr Arg Gln Leu Glu Arg Asp Thr Leu Asn Glu Ala Lys Met Trp 1890 1895 1900	5712
tac atg caa gct ctc cac ctc ttg ggt gac aag cca tac ctc cca ctc Tyr Met Gln Ala Leu His Leu Leu Gly Asp Lys Pro Tyr Leu Pro Leu 1905 1910 1915	5760
agc acc act tgg tcc gac cca agg ttg gac cgt gct gct gac atc acc Ser Thr Thr Trp Ser Asp Pro Arg Leu Asp Arg Ala Ala Asp Ile Thr 1920 1925 1930 1935	5808
act cag aac gct cat gac tct gcc att gtt gct ctc agg cag aac atc Thr Gln Asn Ala His Asp Ser Ala Ile Val Ala Leu Arg Gln Asn Ile 1940 1945 1950	5856
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ttg gct caa agg gtc tac aac ctc aga cac aac ctc tcc att gat ggt Leu Ala Gln Arg Val Tyr Asn Leu Arg His Asn Leu Ser Ile Asp Gly 1985 1990 1995	6000
caa cca ctc tac ctc cca atc tac gcc aca cca gct gac cca aag gct Gln Pro Leu Tyr Leu Pro Ile Tyr Ala Thr Pro Ala Asp Pro Lys Ala 2000 2005 2010 2015	6048
ctt ctc tct gct gtg gct acc agc caa ggt ggt ggc aag ctc cca Leu Leu Ser Ala Ala Val Ala Thr Ser Gln Gly Gly Gly Lys Leu Pro 2020 2025 2030	6096
gag tcc ttc atg tcc ctc tgg agg ttc cca cac atg ttg gag aac gcc Glu Ser Phe Met Ser Leu Trp Arg Phe Pro His Met Leu Glu Asn Ala 2035 2040 2045	6144
cgt ggc atg gtc tcc caa ctc acc cag ttc ggt tcc acc ctc cag aac Arg Gly Met Val Ser Gln Leu Thr Gln Phe Gly Ser Thr Leu Gln Asn 2050 2055 2060	6192
atc att gag agg caa gat gct gag gct ctc aac gct ttg ctc cag aac Ile Ile Glu Arg Gln Asp Ala Glu Ala Leu Asn Ala Leu Leu Gln Asn 2065 2070 2075	6240
cag gca gct gag ttg atc ctc acc aac ttg tcc atc caa gac aag acc Gln Ala Ala Glu Leu Ile Leu Thr Asn Leu Ser Ile Gln Asp Lys Thr 2080 2085 2090 2095	6288
att gag gag ctt gat gct gag aag aca gtc ctt gag aag agc aag gct Ile Glu Glu Leu Asp Ala Glu Lys Thr Val Leu Glu Lys Ser Lys Ala 2100 2105 2110	6336
ggg gcc caa tct cgc ttc gac tcc tac ggc aag ctc tac gat gag aac Gly Ala Gln Ser Arg Phe Asp Ser Tyr Gly Lys Leu Tyr Asp Glu Asn 2115 2120 2125	6384
atc aac gct ggt gag aac cag gcc atg acc ctc agg gct tcc gca gct	6432

Ile Asn Ala Gly Glu Asn Gln Ala Met Thr Leu Arg Ala Ser Ala Ala			
2130	2135	2140	
ggt ctc acc act gct gtc caa gcc tct cgc ttg gct ggt gca gct gct			6480
Gly Leu Thr Thr Ala Val Gln Ala Ser Arg Leu Ala Gly Ala Ala Ala			
2145	2150	2155	
gac ctc gtt cca aac atc ttc ggt ttc gct ggt ggc tcc aga tgg			6528
Asp Leu Val Pro Asn Ile Phe Gly Phe Ala Gly Gly Ser Arg Trp			
2160	2165	2170	2175
ggt gcc att gct gag gct acc ggt tac gtc atg gag ttc tct gcc aac			6576
Gly Ala Ile Ala Glu Ala Thr Gly Tyr Val Met Glu Phe Ser Ala Asn			
2180	2185	2190	
gtc atg aac act gag gct gac aag atc agc caa tct gag acc tac aga			6624
Val Met Asn Thr Glu Ala Asp Lys Ile Ser Gln Ser Glu Thr Tyr Arg			
2195	2200	2205	
agg cgc cgt caa gag tgg gag atc caa agg aac aac gct gag gca gag			6672
Arg Arg Arg Gln Glu Trp Glu Ile Gln Arg Asn Asn Ala Glu Ala Glu			
2210	2215	2220	
ttg aag caa atc gat gct caa ctc aag tcc ttg gct gtc aga agg gag			6720
Leu Lys Gln Ile Asp Ala Gln Leu Lys Ser Leu Ala Val Arg Arg Glu			
2225	2230	2235	
gct gct gtc ctc cag aag acc tcc ctc aag acc caa cag gag caa acc			6768
Ala Ala Val Leu Gln Lys Thr Ser Leu Lys Thr Gln Gln Glu Gln Thr			
2240	2245	2250	2255
cag tcc cag ttg gct ttc ctc caa agg aag ttc tcc aac cag gct ctc			6816
Gln Ser Gln Leu Ala Phe Leu Gln Arg Lys Phe Ser Asn Gln Ala Leu			
2260	2265	2270	
tac aac tgg ctc aga ggc cgc ttg gct gcc atc tac ttc caa ttc tac			6864
Tyr Asn Trp Leu Arg Gly Arg Leu Ala Ala Ile Tyr Phe Gln Phe Tyr			
2275	2280	2285	
gac ctt gct gtg gcc agg tgc ctc atg gct gag caa gcc tac cgc tgg			6912
Asp Leu Ala Val Ala Arg Cys Leu Met Ala Glu Gln Ala Tyr Arg Trp			
2290	2295	2300	
gag ttg aac gat gac tcc gcc agg ttc atc aag cca ggt gct tgg caa			6960
Glu Leu Asn Asp Asp Ser Ala Arg Phe Ile Lys Pro Gly Ala Trp Gln			
2305	2310	2315	
ggc acc tac gct ggt ctc ctt gct ggt gag acc ctc atg ctc tcc ttg			7008
Gly Thr Tyr Ala Gly Leu Leu Ala Gly Glu Thr Leu Met Leu Ser Leu			
2320	2325	2330	2335
gct caa atg gag gat gct cac ctc aag agg gac aag agg gct ttg gag			7056
Ala Gln Met Glu Asp Ala His Leu Lys Arg Asp Lys Arg Ala Leu Glu			
2340	2345	2350	
gtg gag agg aca gtc tcc ctt gct gag gtc tac gct ggt ctc cca aag			7104
Val Glu Arg Thr Val Ser Leu Ala Glu Val Tyr Ala Gly Leu Pro Lys			
2355	2360	2365	
gac aac ggt cca ttc tcc ctt gct caa gag att gac aag ttg gtc agc			7152
Asp Asn Gly Pro Phe Ser Leu Ala Gln Glu Ile Asp Lys Leu Val Ser			

2370	2375	2380	
caa ggt tct ggt tct gct ggt tct ggt aac aac aac ttg gct ttc ggc Gln Gly Ser Gly Ser Ala Gly Ser Gly Asn Asn Asn Leu Ala Phe Gly 2385 2390 2395			7200
gct ggt act gac acc aag acc tcc ctc caa gcc tct gtc tcc ttc gct Ala Gly Thr Asp Thr Lys Thr Ser Leu Gln Ala Ser Val Ser Phe Ala 2400 2405 2410 2415			7248
gac ctc aag atc agg gag gac tac cca gct tcc ctt ggc aag atc agg Asp Leu Lys Ile Arg Glu Asp Tyr Pro Ala Ser Leu Gly Lys Ile Arg 2420 2425 2430			7296
cgc atc aag caa atc tct gtc acc ctc cca gct ctc ttg ggt cca tac Arg Ile Lys Gln Ile Ser Val Thr Leu Pro Ala Leu Leu Gly Pro Tyr 2435 2440 2445			7344
caa gat gtc caa gca atc ctc tcc tac ggt gac aag gct ggt ttg gcg Gln Asp Val Gln Ala Ile Leu Ser Tyr Gly Asp Lys Ala Gly Leu Ala 2450 2455 2460			7392
aac ggt tgc gag gct ctt gct gtc tct cat ggc atg aac gac tct ggt Asn Gly Cys Glu Ala Leu Ala Val Ser His Gly Met Asn Asp Ser Gly 2465 2470 2475			7440
caa ttc caa ctt gac ttc aac gat ggc aag ttc ctc cca ttc gag ggc Gln Phe Gln Leu Asp Phe Asn Asp Gly Lys Phe Leu Pro Phe Glu Gly 2480 2485 2490 2495			7488
att gcc att gac caa ggc acc ctc acc ctc tcc ttc cca aac gct tcc Ile Ala Ile Asp Gln Gly Thr Leu Thr Leu Ser Phe Pro Asn Ala Ser 2500 2505 2510			7536
atg cca gag aag gga aag caa gcc acc atg ctc aag acc ctc aac gat Met Pro Glu Lys Gly Lys Gln Ala Thr Met Leu Lys Thr Leu Asn Asp 2515 2520 2525			7584
atc atc ctc cac atc agg tac acc atc aag tgagctc Ile Ile Leu His Ile Arg Tyr Thr Ile Lys 2530 2535			7621

INTERNATIONAL SEARCH REPORT

Internal Application No
PCT/US 00/22237A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C12N9/52 C12N15/82 C07K14/24 C12N15/11

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

STRAND, EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 08932 A (DOW AGROSCIENCES LLC ;WISCONSIN ALUMNI RES FOUND (US)) 5 March 1998 (1998-03-05) cited in the application SEQ ID NO:11 in this document is the unmodified version of SEQ ID NO:3 of the present application. SEQ ID NO:46 corresponds to SEQ ID NO:5. page 16, line 31 -page 19, line 35 -----	1-7
A	WO 97 13402 A (DOWELANCO) 17 April 1997 (1997-04-17) the whole document -----	1-7

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

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Date of the actual completion of the international search

1 December 2000

Date of mailing of the international search report

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INTERNATIONAL SEARCH REPORT

Information on patent family members

Internal application No

PCT/US 00/22237

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